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FORM PTO-1390 (Modified)  
(REV 11-2000)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

**TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371**

221519US0PCT

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR

**10/089057**

INTERNATIONAL APPLICATION NO.

**PCT/JP00/06913**

INTERNATIONAL FILING DATE

**4 October 2000**

PRIORITY DATE CLAIMED

**4 October 1999 (earliest)**

TITLE OF INVENTION

**GENES FOR HEAT RESISTANT ENZYMES OF AMINO ACID BIOSYNTHETIC PATHWAY DERIVED FROM  
THERMOPHILIC CORYNEFORM BACTERIA**

APPLICANT(S) FOR DO/EO/US

**Seiko HIRANO et al.**

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (24) indicated below.
4. ☒ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
  - a. ☐ is attached hereto (required only if not communicated by the International Bureau).
  - b. ☒ has been communicated by the International Bureau.
  - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
  - a. ☒ is attached hereto.
  - b. ☐ has been previously submitted under 35 U.S.C. 154(d)(4).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
  - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
  - b. ☐ have been communicated by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
  - d. ☒ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
10. ☐ An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).
11. ☐ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☒ A copy of the International Search Report (PCT/ISA/210).

**Items 13 to 20 below concern document(s) or information included:**

13. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☐ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
20. ☐ A second copy of the published international application under 35 U.S.C. 154(d)(4).
21. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
22. ☐ Certificate of Mailing by Express Mail
23. ☒ Other items or information:

**Notice of Priority/ Form PTO-1449****PCT/IB/304/ Drawings (15 sheets)****PCT/IB/308/ Sequence Listing (123 sheets)**

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.492(a)(1)) <div style="font-size: 24pt; font-weight: bold; margin-top: 5px;">10/089057</div>		INTERNATIONAL APPLICATION NO. <div style="font-weight: bold; margin-top: 5px;">PCT/JP00/06913</div>		ATTORNEY'S DOCKET NUMBER <div style="font-weight: bold; margin-top: 5px;">221519US0PCT</div>	
24. The following fees are submitted: <b>BASIC NATIONAL FEE ( 37 CFR 1.492 (a) (1) - (5)) :</b> <div style="margin-top: 5px;"> <input type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO ..... <span style="float: right;">\$1040.00</span> </div> <div style="margin-top: 5px;"> <input checked="" type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO ..... <span style="float: right;">\$890.00</span> </div> <div style="margin-top: 5px;"> <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO ..... <span style="float: right;">\$740.00</span> </div> <div style="margin-top: 5px;"> <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) ..... <span style="float: right;">\$710.00</span> </div> <div style="margin-top: 5px;"> <input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) ..... <span style="float: right;">\$100.00</span> </div> <div style="text-align: right; margin-top: 10px;"> <b>ENTER APPROPRIATE BASIC FEE AMOUNT =</b> </div>				<div style="border: 1px solid black; padding: 5px;"> <b>CALCULATIONS PTO USE ONLY</b>   <div style="border: 1px solid black; height: 100px; margin-top: 5px;"></div> </div>	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).				<div style="border: 1px solid black; padding: 5px;"> <div style="border: 1px solid black; height: 100px; margin-top: 5px;"></div> </div>	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	80 - 20 =	60	x \$18.00	\$1,080.00	
Independent claims	32 - 3 =	29	x \$84.00	\$2,436.00	
Multiple Dependent Claims (check if applicable).				\$280.00	
<b>TOTAL OF ABOVE CALCULATIONS =</b>				\$4,686.00	
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27). The fees indicated above are reduced by 1/2.				\$0.00	
<b>SUBTOTAL =</b>				\$4,686.00	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).				\$0.00	
<b>TOTAL NATIONAL FEE =</b>				\$4,686.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable).				\$0.00	
<b>TOTAL FEES ENCLOSED =</b>				\$4,686.00	
				Amount to be: refunded \$	
				charged \$	
a. <input checked="" type="checkbox"/> A check in the amount of <u>\$4,686.00</u> to cover the above fees is enclosed. b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>15-0030</u> A duplicate copy of this sheet is enclosed. d. <input type="checkbox"/> Fees are to be charged to a credit card. <b>WARNING:</b> Information on this form may become public. <b>Credit card information should not be included on this form.</b> Provide credit card information and authorization on PTO-2038.					
<b>NOTE:</b> Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO:					
<div style="border: 1px solid black; padding: 10px; display: flex; justify-content: space-between; align-items: center;"> <div style="text-align: center;"> <b>Surinder Sachar</b>  <b>Registration No. 34,423</b>    <b>22850</b> </div> <div style="text-align: right;"> <div style="border-bottom: 1px solid black; margin-bottom: 5px;"><i>Surinder Sachar</i></div>           SIGNATURE   <div style="border-bottom: 1px solid black; margin-bottom: 5px;">Norman F. Oblon</div>           NAME   <div style="border-bottom: 1px solid black; margin-bottom: 5px;">24,618</div>           REGISTRATION NUMBER   <div style="border-bottom: 1px solid black; margin-bottom: 5px;">April 3 2002</div>           DATE         </div> </div>					

101 REC'D PCT/PTO 17 DEC 2002 #5  
10/089057

Docket No.221519US0PCT

IN THE UNITED STATES PATENT & TRADEMARK OFFICE

IN RE APPLICATION OF: : ATTN: BOX SEQUENCE

SEIKO HIRANO ET AL :

SERIAL NO. 10/089,057 :

FILED:APRIL 03, 2002 :

FOR:GENES FOR HEAT RESISTANT ENZYMES OF AMINO ACID BIOSYNTHETIC  
PATHWAY DERIVED FROM THERMOPHILIC CORYNEFORM BACTERIA

PRELIMINARY AMENDMENT AND STATEMENT

ASSISTANT COMMISSIONER FOR PATENTS  
WASHINGTON, D.C. 20231

SIR:

Responsive to the Office Communication dated July 17, 2002, Applicants submit a  
substitute Sequence Listing and a corresponding computer-readable Sequence Listing.

IN THE SPECIFICATION

Please amend the specification as follows.

Page 111 (Abstract), after the last line, beginning on a new page, please replace the  
original Sequence Listing with the substitute Sequence Listing attached hereto.

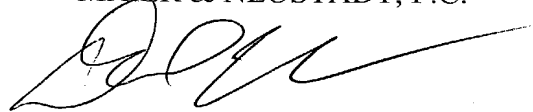
REMARKS

Applicants have now submitted a substitute Sequence Listing and a corresponding computer-readable Sequence Listing. The sequence information recorded in the corresponding computer-readable Sequence Listing is identical to the paper copy of the substitute Sequence Listing. Support for all of the sequences listed in the substitute Sequence Listing is found in the present application as originally filed. No new matter is believed to have been introduced by the submission of the substitute Sequence Listing and the corresponding computer-readable Sequence Listing.

Applicants submit that the present application is ready for examination on the merits. Early notice to this effect is earnestly solicited.

Respectfully submitted,

OBLON, SPIVAK, McCLELLAND,  
MAIER & NEUSTADT, P.C.



Norman F. Oblon  
Attorney of Record  
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Registration No. 45,518



**22850**

(703) 413-3000



## Specification

Genes for Heat resistant Enzymes of Amino Acid  
Biosynthetic Pathway Derived from Thermophilic  
Coryneform Bacteria

5

Technical Field

The present invention relates to heat resistant  
enzyme genes, in particular, genes for enzymes of  
biosynthetic pathway and uptake system of L-amino acids  
such as L-glutamic acid, of *Corynebacterium*  
*thermoaminogenes*, which is a thermophilic coryneform  
bacterium.

15 Background Art

The current main stream of the production of L-  
amino acids such as L-glutamic acid is the fermentative  
production utilizing coryneform bacteria. As for the  
fermentative production of L-amino acids, it has been  
attempted to reduce the cost based on breeding of  
strains with superior productivity and development of  
fermentation techniques. Although conventional attempts  
for realizing the cost reduction were mainly directed to  
achieving higher yield, energy required for cooling the  
fermentation heat generated during the culture cannot be  
ignored in addition to the raw material as the factors  
concerning the fermentation cost. That is, as for usual

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microorganisms used for the fermentation, the temperature of the medium rises due to fermentation heat generated by the microorganism themselves during the fermentation, and hence enzymes required for the fermentation may be inactivated or the productive bacteria may be killed. Therefore, it is necessary to cool the medium during the fermentation. Accordingly, in order to reduce the cooling cost, fermentation at high temperatures has been studied for many years. Moreover, if high temperature fermentation becomes possible, the reaction rate may also be improved. However, as for the L-amino acid fermentation, effective high temperature culture has not been realized so far.

*Corynebacterium thermoaminogenes* is a bacterium classified into coryneform bacteria like *Corynebacterium glutamicum* (*Brevibacterium lactofermentum*), which is commonly used for the fermentation of L-amino acids. However, it shows the optimum growth temperature of 37-43°C, which is higher than that of *Corynebacterium glutamicum*, i.e., 30-35°C, and shows the optimum temperature for L-glutamic acid production of 42-45°C, which is considerably shifted to the high temperature region (Japanese Patent Laid-open (Kokai) No. 63-240779/1988).

Meanwhile, there have been developed techniques for enhancing L-amino acid producing ability of *Corynebacterium* and *Brevibacterium* bacteria by

introducing a gene coding for an L-amino acid synthesis system enzyme derived from *Escherichia coli* or *Corynebacterium glutamicum* into them. Examples of such an enzyme include, for example, citrate synthase  
5 (Japanese Patent Publication (Kokoku) No. 7-121228/1995), which is an enzyme of the L-glutamic acid biosynthetic pathway, glutamate dehydrogenase (Japanese Patent Laid-open No. 61-268185/1986), isocitrate dehydrogenase, aconitate hydratase (Japanese Patent Laid-open No. 63-  
10 214189) and so forth.

However, any L-amino acid biosynthesis enzymes and genes coding for them derived from thermophilic coryneform bacteria have not been reported.

15 Disclosure of the Invention

An object of the present invention is to provide genes coding for enzymes derived from *Corynebacterium thermoaminogenes*, preferably enzymes that function at a temperature higher than those of *Corynebacterium*  
20 *glutamicum*.

The inventors of the present invention extensively studied in order to achieve the aforementioned object. As a result, they successfully isolated genes coding for enzymes of the amino acid biosynthetic pathway of  
25 *Corynebacterium thermoaminogenes*, or genes coding for proteins involved in the uptake of amino acids into cells, and thus achieved the present invention.

That is, the present invention provides the followings.

- 5 (1) A protein having the amino acid sequence of SEQ ID NO: 2 or the amino acid sequence of SEQ ID NO: 2 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has isocitrate lyase activity and shows 30% or more of residual activity after a heat treatment at 50°C for 5 minutes.
- 10 (2) A protein having the amino acid sequence of SEQ ID NO: 4 or the amino acid sequence of SEQ ID NO: 4 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which is involved in acyl Co-A carboxylase activity derived  
15 from *Corynebacterium thermoaminogenes*.
- (3) A protein having the amino acid sequence of SEQ ID NO: 6 or the amino acid sequence of SEQ ID NO: 6 including substitution, deletion, insertion, addition or  
20 inversion of one or several amino acids residues, which has DtsR activity derived from *Corynebacterium thermoaminogenes*.
- (4) A protein having the amino acid sequence of SEQ ID NO: 8 or the amino acid sequence of SEQ ID NO: 8 including substitution, deletion, insertion, addition or  
25 inversion of one or several amino acids residues, which has DtsR activity derived from *Corynebacterium thermoaminogenes*.

- (5) A protein having the amino acid sequence of SEQ ID NO: 10 or the amino acid sequence of SEQ ID NO: 10 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which shows phosphofructokinase activity at 60°C in an equivalent or higher degree compared with the activity at 30°C.
- (6) A protein having the amino acid sequence of SEQ ID NO: 94 or the amino acid sequence of SEQ ID NO: 94 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has activity for imparting sucrose assimilating ability to *Corynebacterium thermoaminogenes*.
- (7) A protein having any one of the amino acid sequences of SEQ ID NOS: 17-20 or the amino acid sequence of any one of SEQ ID NOS: 17-20 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has a function involved in glutamic acid uptake and derived from *Corynebacterium thermoaminogenes*.
- (8) A protein having the amino acid sequence of SEQ ID NO: 22 or the amino acid sequence of SEQ ID NO: 22 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has pyruvate dehydrogenase activity derived from *Corynebacterium thermoaminogenes*.
- (9) A protein having the amino acid sequence of SEQ ID

NO: 24 or the amino acid sequence of SEQ ID NO: 24  
including substitution, deletion, insertion, addition or  
inversion of one or several amino acids residues, which  
has pyruvate carboxylase activity derived from

5 *Corynebacterium thermoaminogenes*.

(10) A protein having the amino acid sequence of SEQ ID  
NO: 26 or the amino acid sequence of SEQ ID NO: 26  
including substitution, deletion, insertion, addition or  
inversion of one or several amino acids residues, which  
10 has phosphoenolpyruvate carboxylase activity and shows  
50% or more of residual activity after a heat treatment  
at 45°C for 5 minutes.

(11) A protein having the amino acid sequence of SEQ ID  
NO: 28 or the amino acid sequence of SEQ ID NO: 28  
15 including substitution, deletion, insertion, addition or  
inversion of one or several amino acids residues, which  
has aconitase activity and shows 30% or more of residual  
activity after a heat treatment at 50°C for 3 minutes.

(12) A protein having the amino acid sequence of SEQ ID  
20 NO: 30 or the amino acid sequence of SEQ ID NO: 30  
including substitution, deletion, insertion, addition or  
inversion of one or several amino acids residues, which  
has isocitrate dehydrogenase activity and shows 50% or  
more of residual activity after a heat treatment at 45°C  
25 for 10 minutes.

(13) A protein having the amino acid sequence of SEQ ID  
NO: 32 or the amino acid sequence of SEQ ID NO: 32

including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has dihydrolipoamide dehydrogenase activity derived from *Corynebacterium thermoaminogenes*.

5 (14) A protein having the amino acid sequence of SEQ ID NO: 34 or the amino acid sequence of SEQ ID NO: 34 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has 2-oxoglutarate dehydrogenase activity and shows 30%  
10 or more of residual activity after a heat treatment at 50°C for 10 minutes.

(15) A protein having the amino acid sequence of SEQ ID NO: 80 in Sequence Listing or the amino acid sequence of SEQ ID NO: 80 including substitution, deletion,  
15 insertion, addition or inversion of one or several amino acids residues, which shows glutamate dehydrogenase activity at 42°C in an equivalent or higher degree compared with the activity at 37°C.

(16) A protein having the amino acid sequence of SEQ ID  
20 NO: 90 in Sequence Listing or the amino acid sequence of SEQ ID NO: 90 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which shows citrate synthase activity at 37°C in an equivalent or higher degree compared with the  
25 activity at 23°C.

(17) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 2 or the amino acid sequence

of SEQ ID NO: 2 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having isocitrate lyase activity.

(18) The DNA according to (17), which is a DNA defined  
5 in the following (a1) or (b1):

(a1) a DNA which comprises the nucleotide sequence of SEQ ID NO: 1 in Sequence Listing,

(b1) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 1 in Sequence Listing  
10 or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having isocitrate lyase activity.

(19) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 4 or the amino acid sequence  
15 of SEQ ID NO: 4 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and involved in acyl Co-A carboxylase activity.

(20) The DNA according to (19), which is a DNA defined  
20 in the following (a2) or (b2):

(a2) a DNA which comprises the nucleotide sequence of SEQ ID NO: 3 in Sequence Listing,

(b2) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 3 in Sequence Listing  
25 or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein involved in acyl Co-A carboxylase activity.



(21) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 6 or the amino acid sequence of SEQ ID NO: 6 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having DtsR activity.

(22) The DNA according to (21), which is a DNA defined in the following (a3) or (b3):

(a3) a DNA which comprises the nucleotide sequence of SEQ ID NO: 5 in Sequence Listing,

(b3) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 5 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having DtsR activity.

(23) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 8 or the amino acid sequence of SEQ ID NO: 8 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having DtsR activity.

(24) The DNA according to (23), which is a DNA defined in the following (a4) or (b4):

(a4) a DNA which comprises the nucleotide sequence of SEQ ID NO: 7 in Sequence Listing,

(b4) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 7 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein

having DtsR activity.

(25) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 10 or the amino acid sequence of SEQ ID NO: 10 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having phosphofructokinase activity.

(26) The DNA according to (25), which is a DNA defined in the following (a5) or (b5):

10 (a5) a DNA which comprises the nucleotide sequence of SEQ ID NO: 9 in Sequence Listing,

(b5) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 9 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having phosphofructokinase activity.

15 (27) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 93 or the amino acid sequence of SEQ ID NO: 93 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having invertase activity.

(28) The DNA according to (27), which is a DNA defined in the following (a6) or (b6):

25 (a6) a DNA which comprises the nucleotide sequence of SEQ ID NO: 93 in Sequence Listing,

(b6) a DNA which is hybridizable with the

nucleotide sequence of SEQ ID NO: 93 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having invertase activity.

- 5 (29) A DNA which codes for a protein having any one of the amino acid sequences of SEQ ID NOS: 17-20 or the amino acid sequence of any one of SEQ ID NOS: 17-20 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and  
10 having a function involved in glutamic acid uptake.  
(30) The DNA according to (29), which is a DNA defined in the following (a7) or (b7):

(a7) a DNA which comprises the nucleotide sequence of SEQ ID NO: 16 in Sequence Listing,

- 15 (b7) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 16 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having a function involved in glutamic acid uptake.

- 20 (31) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 22 or the amino acid sequence of SEQ ID NO: 22 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having pyruvate  
25 dehydrogenase activity.

(32) The DNA according to (31), which is a DNA defined in the following (a8) or (b8):

(a8) a DNA which comprises the nucleotide sequence of SEQ ID NO: 21 in Sequence Listing,

(b8) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 21 in Sequence Listing  
5 or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having pyruvate dehydrogenase activity.

(33) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 24 or the amino acid  
10 sequence of SEQ ID NO: 24 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having pyruvate carboxylase activity.

(34) A DNA according to (33), which is a DNA defined in  
15 the following (a9) or (b9):

(a9) a DNA which comprises the nucleotide sequence of SEQ ID NO: 23 in Sequence Listing,

(b9) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 23 in Sequence Listing  
20 or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having pyruvate carboxylase activity.

(35) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 26 or the amino acid  
25 sequence of SEQ ID NO: 26 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having

phosphoenolpyruvate carboxylase activity.

(36) The DNA according to (35), which is a DNA defined in the following (a10) or (b10):

5 (a10) a DNA which comprises the nucleotide sequence of SEQ ID NO: 25 in Sequence Listing,

(b10) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 25 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein  
10 having phosphoenolpyruvate carboxylase activity.

(37) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 28 or the amino acid sequence of SEQ ID NO: 28 including substitution, deletion, insertion, addition or inversion of one or  
15 several amino acids residues, and having aconitase activity.

(38) The DNA according to (37), which is a DNA defined in the following (a11) or (b11):

(a11) a DNA which comprises the nucleotide  
20 sequence of SEQ ID NO: 27 in Sequence Listing,

(b11) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein  
25 having aconitase activity.

(39) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 30 or the amino acid

sequence of SEQ ID NO: 30 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having isocitrate dehydrogenase activity.

- 5 (40) The DNA according to (39), which is a DNA defined in the following (a12) or (b12):

(a12) a DNA which comprises the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing,

- 10 (b12) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having isocitrate dehydrogenase activity.

- 15 (41) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 32 or the amino acid sequence of SEQ ID NO: 32 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having dihydrolipoamide dehydrogenase activity.

- 20 (42) The DNA according to (41), which is a DNA defined in the following (a13) or (b13):

(a13) a DNA which comprises the nucleotide sequence of SEQ ID NO: 31 in Sequence Listing,

- 25 (b13) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 31 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein

having dihydrolipoamide dehydrogenase activity.

(43) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 34 or the amino acid sequence of SEQ ID NO: 34 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having 2-oxoglutarate dehydrogenase activity.

(44) The DNA according to (43), which is a DNA defined in the following (a14) or (b14):

10 (a14) a DNA which comprises the nucleotide sequence of SEQ ID NO: 33 in Sequence Listing,

(b14) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 33 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having 2-oxoglutarate dehydrogenase activity.

15 (45) A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 80 in Sequence Listing or the amino acid sequence of SEQ ID NO: 80 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and showing glutamate dehydrogenase activity at 42°C in an equivalent or higher degree compared with the activity at 37°C.

25 (46) The DNA according to (45), which is a DNA defined in the following (a15) or (b15):

(a15) a DNA which comprises the nucleotide

sequence of SEQ ID NO: 79 in Sequence Listing,

(b15) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 79 in Sequence Listing or a primer prepared based on the nucleotide sequence  
5 under a stringent condition, and codes for a protein showing glutamate dehydrogenase activity at 42°C in an equivalent or higher degree compared with the activity at 37°C.

(47) A DNA which codes for a protein having the amino  
10 acid sequence of SEQ ID NO: 90 in Sequence Listing or the amino acid sequence of SEQ ID NO: 90 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and showing citrate synthase activity at 37°C in an equivalent or  
15 higher degree compared with the activity at 23°C.

(48) The DNA according to (47), which is a DNA defined in the following (a16) or (b16):

(a16) a DNA which comprises the nucleotide sequence of SEQ ID NO: 89 in Sequence Listing,

20 (b16) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 89 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein showing citrate synthase activity at 37°C in an  
25 equivalent or higher degree compared with the activity at 23°C.

(49) A method for producing L-amino acid, which



comprises culturing a microorganism introduced with a DNA according to any one of (17) to (48) in a medium to produce and accumulate L-amino acid in the medium, and collecting the L-amino acid from the medium.

5           The term "DNA of the present invention" is used hereinafter for referring to either one or all of the aforementioned DNAs.

Hereafter, the present invention will be explained in detail.

10           The nucleotide sequences of the DNA of the present invention, names of the genes, and the proteins encoded by the DNA of the present invention are shown in Table 1.

Table 1

Nucleotide sequence	Name of gene	Encoded protein (abbreviation)
SEQ ID NO: 1	<i>aceA</i>	Isocitrate lyase (ICL)
SEQ ID NO: 3	<i>accBC</i>	acyl Co-A carboxylase BC subunit
SEQ ID NO: 5	<i>dtsR1</i>	DTSR1 protein
SEQ ID NO: 7	<i>dtsR2</i>	DTSR2 protein
SEQ ID NO: 9	<i>pfk</i>	Phosphofructokinase
SEQ ID NOS: 11, 13, 15, 93	<i>scrB</i>	Invertase
SEQ ID NO: 16	<i>gluABCD</i>	glutamic acid uptake system
SEQ ID NO: 21	<i>pdhA</i>	pyruvate dehydrogenase
SEQ ID NO: 23	<i>pc</i>	pyruvate carboxylase
SEQ ID NO: 25	<i>ppc</i>	phosphoenolpyruvate carboxylase
SEQ ID NO: 27	<i>acn</i>	aconitase
SEQ ID NO: 29	<i>icd</i>	isocitrate dehydrogenase
SEQ ID NO: 31	<i>lpd</i>	dihydrolipoamide dehydrogenase
SEQ ID NO: 33	<i>odhA</i>	2-oxoglutarate dehydrogenase
SEQ ID NO: 79	<i>gdh</i>	glutamate dehydrogenase
SEQ ID NO: 89	<i>glta</i>	citrate synthase

The open reading frames (ORFs) of SEQ ID NOS: 3, 23, 25, 31 and 33 and the fourth ORF of SEQ ID NO: 16 all start from GTG. Although the amino acids encoded by these GTG are indicated as valine in Sequence Listing, they may be methionine.

The sequence of SEQ ID NO: 16 contains four ORFs, which correspond to *gluA*, *gluB*, *gluC* and *gluD* in this order from the 5' end side.

The aforementioned DNA sequences were isolated from chromosomal DNA of the *Corynebacterium thermoaminogenes* AJ12310 strain (FERM BP-1542). However, the DNA sequences shown in SEQ ID NOS: 11 and 13 were isolated from *Corynebacterium thermoaminogenes* AJ12340 strain (FERM BP-1539) and AJ12309 strain (FERM BP-1541),

respectively, which had invertase activity and sucrose assimilating property, because the AJ12310 strain did not have invertase activity and sucrose assimilating property, and the *scrB* gene isolated from the strain had  
5 not any open reading frame.

The *Corynebacterium thermoaminogenes* AJ12310 strain (also referred to as YS-314 strain) and AJ12309 strain (also referred to as YS-155 strain) were deposited at the National Institute of Bioscience and  
10 Human-Technology, Agency of Industrial Science and Technology, Ministry of International Trade and Industry (postal code: 305-8566, 1-3, Higashi 1-chome, Tsukuba-shi, Ibaraki-ken, Japan) on March 13, 1987 and given deposition numbers of FERM P-9246 and FERM P-9245,  
15 respectively. Then, they were transferred to international depositions under the provisions of the Budapest Treaty on October 27, 1987, and given deposition numbers of FERM BP-1542 and FERM BP-1541, respectively.

20 The AJ12340 strain (also referred to as YS-40 strain) was deposited at the National Institute of Bioscience and Human-Technology, Agency of Industrial Science and Technology, Ministry of International Trade and Industry (postal code: 305-8566, 1-3, Higashi 1-  
25 chome, Tsukuba-shi, Ibaraki-ken, Japan) on March 10, 1987 and given a deposition number of FERM P-9277. Then, it was transferred to an international deposition under

the provisions of the Budapest Treaty on October 27, 1987, and given a deposition number of FERM BP-1539.

The nucleotide sequences shown in SEQ ID NOS: 11, 13 and 15 are partial sequences of *scrB*, and the  
5 sequences of SEQ ID NOS: 11 and 13 code for partial amino acid sequences of invertase shown in SEQ ID NOS: 12 and 14.

A DNA sequence containing a partial fragment of a target gene can be obtained by comparing already  
10 reported nucleotide sequences for the target gene of various microorganisms such as *Brevibacterium lactofermentum* to select a region containing a well-conserved nucleotide sequence, and carrying out PCR using primers designed based on the nucleotide sequence  
15 of the region and chromosomal DNA of *Corynebacterium thermoaminogenes* as a template. Further, by performing hybridization using the obtained DNA fragment or a probe prepared based on the sequence of the fragment to screen a chromosomal DNA library of *Corynebacterium*  
20 *thermoaminogenes*, a DNA fragment containing the gene in its full length can be obtained. A DNA fragment containing the gene in its full length can also be obtained by performing genome walking using the obtained partial fragment of the gene. The genome walking can be  
25 carried out by using a commercially available kit, for example, TaKaRa LA PCR in vitro Cloning Kit (produced by Takara Shuzo).

For example, a partial sequence of DNA coding for glutamate dehydrogenase (henceforth the DNA is also referred to as "*gdh*", and the enzyme is also referred to as "GDH") can be obtained from chromosomal DNA of

- 5 *Corynebacterium thermoaminogenes* such as the *Corynebacterium thermoaminogenes* AJ12310 strain by PCR (polymerase chain reaction) using the chromosomal DNA as a template and primers having the nucleotide sequences shown in SEQ ID NOS: 77 and 78 of Sequence Listing.
- 10 Further, by performing genome walking using the obtained partial fragment, the whole *gdh* gene can be obtained.

- Further, a partial sequence of DNA coding for citrate synthase (henceforth the DNA is also referred to as "*gltA*", and the enzyme is also referred to as "CS")
- 15 can be obtained from chromosomal DNA of *Corynebacterium thermoaminogenes* such as the *Corynebacterium thermoaminogenes* AJ12310 strain by PCR (polymerase chain reaction) using the chromosomal DNA as a template and primers having the nucleotide sequences shown in SEQ ID
- 20 NOS: 83 and 84 of Sequence Listing. Further, by performing genome walking using the obtained partial fragment, the whole *gltA* gene can be obtained.

- The nucleotide sequences of the aforementioned primers were designed based on a nucleotide sequence in
- 25 a region containing a well-conserved nucleotide sequence among the already reported *gdh* genes or *gltA* genes of various microorganisms, which region was found by

comparison of the genes.

As for DNA sequences coding for the other enzymes, partial fragments coding for those enzymes can be similarly obtained by using the primers mentioned in  
5 Table 1, and the genes in full length can be obtained by using the obtained partial fragments.

While the DNA of the present invention was obtained as described above, it can also be obtained from a chromosomal DNA library of *Corynebacterium*  
10 *thermoaminogenes* by hybridization using an oligonucleotide prepared based on the nucleotide sequences of the DNA of the present invention as a probe.

Methods for preparation of chromosomal DNA, construction of chromosomal DNA library, hybridization,  
15 PCR, preparation of plasmid DNA, digestion and ligation of DNA, transformation and so forth are described in Sambrook, J., Fritsch, E.F., Maniatis, T., Molecular Cloning, Cold Spring Harbor Laboratory Press, 1.21 (1989). Further, genome walking can be performed by  
20 using a commercially available kit, for example, TaKaRa LA PCR in vitro Cloning Kit (produced by Takara Shuzo).

Specific methods for obtaining the DNA of the present invention will be explained hereafter.

First, chromosomal DNA of *Corynebacterium*  
25 *thermoaminogenes* is digested with a suitable restriction enzyme, for example, *Sau3AI*, and fractionated by agarose gel electrophoresis to obtain a DNA fragment of about 4

to 6 kb. The obtained DNA fragment is inserted into a cloning vector such as pHSG399, and *Escherichia coli* is transformed with the obtained recombinant plasmid to produce a plasmid library of the chromosomal DNA.

5           Separately, primers are produced for use in selecting a clone containing a target gene from a plasmid library by PCR. These primers are designed based on conserved amino acid regions from various microorganisms corresponding to the gene of interest.

10          In the design of primers, a plurality of primer sets are designed considering the codon usage of coryneform bacteria.

          Then, in order to investigate propriety of the produced primers, PCR is performed by using these  
15          primers and chromosomal DNA of *Corynebacterium thermoaminogenes* as a template. Further, PCR is performed by using primers from which an amplification fragment has been obtained as primers for screening and a recombinant plasmid prepared from the plasmid library  
20          as a template to select a clone containing the target DNA fragment. This operation can be quickly carried out by performing the PCR for every batch including several tens of transformant strains as primary screening and performing colony PCR for the batch with which an  
25          amplification fragment was obtained as secondary screening. The fragment lengths of the amplified genes

are shown in Tables 2 to 7.

If a transformant selected as described above contains a target gene is confirmed by preparing a recombinant DNA from the transformant selected as described above, determining the nucleotide sequence of the inserted fragment by the dideoxy termination method, and comparing the nucleotide sequence with a known gene sequence.

When the obtained DNA fragment contains a part of the target gene, the deleted part is obtained by genome walking.

The DNA of the present invention may code for a protein including substitution, deletion, insertion, addition or inversion of one or several amino acid residues, so long as the encoded protein has its original function. The number meant by the term "several" may vary depending on positions in the three-dimensional structure of protein or kinds of amino acid residues. However, in general, such a protein preferably shows homology of 30 to 40% or more, more preferably 55 to 65% or more, with respect to a corresponding whole amino acid sequence of the protein. More specifically, the term "several" means a number of 2 to several hundreds, preferably 2 to several tens, more preferably 2 to 10.

Nucleotide and amino acid sequence were analyzed by, for example, the method developed by Lipman and



Peason (Science, 227, 1435-1441, 1985) by using commercially available software such as Genetyx-Mac computer program (Software Development Co., Tokyo, Japan).

5           GDH may be one showing homology of 40 to 80% or more, preferably 80 to 90% or more, for the total amino acid sequence constituting GDH, and showing GDH activity at 42°C equivalent to or higher than the activity at 37°C. In this case, the term "several" means a number  
10 of 2 to 30, preferably 2 to 50, more preferably 2 to 10.

          CS may be one showing homology of 40 to 80% or more, preferably 80 to 90% or more, for the total amino acid sequence constituting CS, and showing CS activity at 37°C equivalent to or higher than the activity at  
15 23°C. In this case, the term "several" means a number of 2 to 300, preferably 2 to 50, more preferably 2 to 10.

          A DNA, which codes for the substantially same protein as the original protein as described above, can be obtained by, for example, modifying the nucleotide  
20 sequence, for example, by means of the site-directed mutagenesis so that one or more amino acid residues at a specific site should involve substitution, deletion, insertion, addition or inversion. A DNA modified as described above may also be obtained by a conventionally  
25 known mutation treatment. The mutation treatment includes a method for treating DNA coding for a target gene in vitro, for example, with hydroxylamine, and a

method for treating a microorganism, for example, a bacterium belonging to the genus *Escherichia*, harboring DNA coding for the target gene with ultraviolet irradiation or a mutating agent usually used for the mutation treatment such as N-methyl-N'-nitro-N-nitrosoguanidine (NTG) and nitrous acid.

The substitution, deletion, insertion, addition, or inversion of nucleotides as described above also includes mutant or variant that naturally occurs due to the difference of strains of *Corynebacterium thermoaminogenes* or the like.

A DNA coding for substantially the same protein as the original protein can be obtained by expressing DNA having a mutation in an appropriate cell, and investigating activity or function of the expressed product protein. The DNA coding for substantially the same protein as the original protein can also be obtained by, for example, isolating a DNA which is hybridizable with a DNA having each of the nucleotide sequences of the sequences of which sequence numbers are mentioned in Table 1 or a coding region thereof, or a probe designed based on the nucleotide sequence under a stringent condition, and which codes for a protein having the activity originally possessed by the protein, from DNA coding for a protein having a mutation or from a cell harboring it. The activity preferably means each enzymatic activity at 42°C for GDH or 37°C for CS.

The aforementioned probe can be prepared from a DNA having any one of the nucleotide sequences of which sequence numbers are shown in Table 1 or a DNA having any one of the nucleotide sequences by PCR using  
5 suitable primers.

The "stringent condition" referred to herein is a condition under which so-called specific hybrid is formed, and non-specific hybrid is not formed. It is difficult to clearly express this condition by using any  
10 numerical value. However, for example, the stringent condition includes a condition under which DNAs having high homology, for example, DNAs having homology of not less than 50% are hybridized with each other, and DNAs having homology lower than the above are not hybridized  
15 with each other. Alternatively, the stringent condition is exemplified by a condition under which DNAs are hybridized with each other at a salt concentration corresponding to an ordinary condition of washing in Southern hybridization, *i.e.*, 60°C, 1 x SSC, 0.1% SDS,  
20 preferably 0.1 x SSC, 0.1% SDS.

The gene, which is hybridizable under the condition as described above, includes those having a stop codon generated in the gene, and those having no activity due to mutation of active site. However, such  
25 genes can be easily removed by ligating the genes with a commercially available activity expression vector, and measuring the activity or function.

A protein corresponding to each DNA of the present invention can be produced by expressing the DNA in a suitable host-vector system.

As the host used for the expression of a gene,  
5 there can be mentioned various prokaryotic cells including *Brevibacterium lactofermentum* (*Corynebacterium glutamicum*), coryneform bacteria such as *Corynebacterium thermoaminogenes*, *Escherichia coli*, *Bacillus subtilis* and so forth, and various eucaryotic cells including  
10 *Saccharomyces cerevisiae*, animal cells and plant cells. Among these, prokaryotic cells, in particular, coryneform bacteria and *Escherichia coli* are preferred.

If the DNA of the present invention is ligated to a vector DNA autonomously replicable in cells of  
15 *Escherichia coli* and/or coryneform bacteria and so forth to form a recombinant DNA, and this recombinant DNA is introduced into an *Escherichia coli* cell, the subsequent procedure becomes easy. The vector autonomously replicable in *Escherichia coli* cells is preferably a  
20 plasmid vector autonomously replicable in the host cell, and examples thereof include pUC19, pUC18, pBR322, pHSG299, pHSG399, pHSG398, RSF1010 and so forth.

As the vector autonomously replicable in coryneform bacterium cells, there can be mentioned  
25 pAM330 (refer to Japanese Patent Laid-open No. 58-67699/1983), pHM1519 (refer to Japanese Patent Laid-open No. 58-77895/1983) and so forth. Moreover, if a DNA

fragment having an ability to make a plasmid  
autonomously replicable in coryneform bacteria is taken  
out from these vectors and inserted into the  
aforementioned vectors for *Escherichia coli*, they can be  
5 used as a so-called shuttle vector autonomously  
replicable in both of *Escherichia coli* and coryneform  
bacteria.

Examples of such a shuttle vector include those  
mentioned below. There are also indicated  
10 microorganisms that harbor each vector, and accession  
numbers thereof at international depositories are shown  
in the parentheses, respectively.

pAJ655 *Escherichia coli* AJ11882 (FERM BP-136)  
*Corynebacterium glutamicum* SR8201 (ATCC39135)  
15 pAJ1844 *Escherichia coli* AJ11883 (FERM BP-137)  
*Corynebacterium glutamicum* SR8202 (ATCC39136)  
pAJ611 *Escherichia coli* AJ11884 (FERM BP-138)  
pAJ3148 *Corynebacterium glutamicum* SR8203 (ATCC39137)  
pAJ440 *Bacillus subtilis* AJ11901 (FERM BP-140)  
20 pH4 *Escherichia coli* AJ12617 (FERM BP-3532)

In order to prepare a recombinant DNA by ligating  
the DNA of the present invention and a vector that  
functions in coryneform bacteria, the vector is digested  
25 with a restriction enzyme that provides an end  
corresponding to an end of the DNA of the present  
invention. The ligation is normally attained by using a

ligase such as T4 DNA ligase.

To introduce the recombinant DNA prepared as described above into a host such as coryneform bacteria, any known transformation methods that have hitherto been reported can be employed. For instance, employable are a method of treating recipient cells with calcium chloride so as to increase the permeability for DNA, which has been reported for *Escherichia coli* K-12 (Mandel, M. and Higa, A., *J. Mol. Biol.*, 53, 159 (1970)), and a method of preparing competent cells from cells which are at the growth phase followed by introducing the DNA thereinto, which has been reported for *Bacillus subtilis* (Duncan, C.H., Wilson, G.A. and Young, F.E., *Gene*, 1, 153 (1977)). In addition to these, also employable is a method of making DNA-recipient cells into protoplasts or spheroplasts, which can easily take up recombinant DNA, followed by introducing the recombinant DNA into the cells, which is known to be applicable to *Bacillus subtilis*, actinomycetes and yeasts (Chang, S. and Choen, S.N., *Molec. Gen. Genet.*, 168, 111 (1979); Bibb, M.J., Ward, J.M. and Hopwood, O.A., *Nature*, 274, 398 (1978); Hinnen, A., Hicks, J.B. and Fink, G.R., *Proc. Natl. Sci. USA*, 75, 1929 (1978)). The transformation of coryneform bacteria can be effectively performed by the electric pulse method (refer to Japanese Patent Laid-open No. 2-207791).

As for the transformation of thermophilic

coryneform bacteria such as *Corynebacterium thermoaminogenes*, it can be efficiently performed by treating cells with an agent that changes the structure of cell walls of the host cells, and applying an electric pulse to a solution containing DNA and the cells of which structure of the cell walls have been changed. The aforementioned agent is an agent that can change the structure of cell walls so that the cells can uptake the DNA when an electric pulse is applied to a solution containing the cells treated with the agent and the DNA (henceforth also referred to as a "cell wall treatment agent"). Examples of such an agent include agents that inhibit normal synthesis of bacterial cell wall and agents that lyse bacterial cell walls. Specific examples thereof include lysozyme, penicillin G, glycine and so forth.

Those cell wall treatment agents may be used each alone, or two or more kinds of them may be used in combination. Among the aforementioned agents, lysozyme and penicillin G are preferred, and lysozyme is particularly preferred.

Furthermore, the transformation of *Corynebacterium thermoaminogenes* can also be performed by applying an electric pulse to a solution containing DNA and the host cells of which cell walls has been weakened by a physical method such as ultrasonication (*FEMS Microbiology Letters*, 151, 135-138 (1987)).

In order to efficiently express a gene contained in the DNA of the present invention, a promoter that functions in the host cell such as lac, trp and  $P_L$  may be ligated upstream from the coding region of the gene.

5 If a vector containing a promoter is used as the vector, ligation of each gene, vector and promoter can be attained by one step.

The proteins of the present invention, which can be produced as described above, can be purified as  
10 required from a cell extract or medium by using usual methods for purifying enzymes such as ion exchange chromatography, gel filtration chromatography, adsorption chromatography, salting out and solvent precipitation.

15 It is expected that the proteins of the present invention are excellent in thermal stability or exhibit higher activity at high temperatures compared with the corresponding proteins of *Corynebacterium glutamicum* and so forth. For example, GDH of *Brevibacterium*  
20 *lactofermentum* shows the highest GDH specific activity around 37°C, and the activity is markedly reduced around 42°C. However, GDH of the present invention shows at 42°C the GDH activity equivalent to or higher than the activity at 37°C. In a preferred embodiment, GDH of the  
25 present invention shows the highest specific activity around 42°C, and shows the activity even at 45°C.

The GDH activity can be measured by, for example,



adding the enzyme to 100 mM Tris-HCl (pH 8.0), 20 mM  $\text{NH}_4\text{Cl}$ , 10 mM sodium  $\alpha$ -ketoglutarate, 0.25 mM NADPH, and determining change of absorbance at 340 nm (Molecular Microbiology 6, 317-326 (1992)).

5           Further, CS of *Brevibacterium lactofermentum* shows the highest CS specific activity around 23°C, and the activity is markedly reduced around 33°C. To the contrary, CS of the present invention shows at 37°C the CS activity equivalent to or higher than the activity at  
10   23°C. In a preferred embodiment, CS of the present invention shows reaction temperature-dependently higher activity up to around 37°C, and shows, even at 40°C, about 40% of the activity with respect to the activity at 37°C.

15           The CS activity can be measured by, for example, the method described in Methods in Enzymol., 13, 3-11 (1969).

          Further, other proteins of the present invention typically have the following characteristics. The  
20   isocitrate lyase has 30% or more of residual activity after a heat treatment at 50°C for 5 minutes. The phosphofructokinase has, at 60°C, the activity equivalent to or higher than the activity at 30°C. The phosphoenolpyruvate carboxylase has 50% or more of  
25   residual activity after a heat treatment at 45°C for 5 minutes. The aconitase has 30% or more of residual activity after a heat treatment at 50°C for 3 minutes.

The isocitrate dehydrogenase has 50% or more of residual activity after a heat treatment at 45°C for 10 minutes.

The 2-oxoglutarate dehydrogenase has 30% or more of residual activity after a heat treatment at 50°C for 10  
5 minutes.

The proteins of the present invention can also be obtained from cell extracts of *Corynebacterium thermoaminogenes* such as the *Corynebacterium thermoaminogenes* AJ12310 strain by using each activity  
10 as an index and usual purification methods for purifying enzymes such as ion exchange chromatography, gel filtration chromatography, adsorption chromatography, salting out and solvent precipitation.

Among the DNA of the present invention, *pfk*, *pdhA*,  
15 *pc*, *ppc*, *acn*, *icd*, *gdh* and *gltA* (names of the enzymes encoded by these are shown in Table 1) can be introduced into L-amino acid production bacteria such as coryneform bacteria to enhance their L-amino acid producing ability. It is also expected that coryneform bacteria introduced  
20 with the DNA of the present invention become possible to produce L-amino acid at a temperature higher than usual. The L-amino acid includes L-glutamic acid, L-aspartic acid, L-lysine, L-arginine, L-proline, L-glutamine and so forth.

25 For example, it is expected that L-glutamic acid production bacteria such as coryneform bacteria

introduced with the *gdh* gene or *gltA* gene come to be able to produce L-glutamic acid at a temperature higher than usual. Further, although CS of *Brevibacterium lactofermentum* may not fully function at a usual culture temperature, for example, 31.5°C, the activity can be enhanced by introducing the *gltA* gene of the present invention.

Further, *dtsR1* and *dtsR2* are genes that code for proteins imparting resistance to surfactant to coryneform bacteria (DTSR protein), and coryneform L-glutamic acid producing bacteria of which these genes are disrupted produce a marked amount of L-glutamic acid even under a condition where biotin is present in such an amount that a wild strain becomes to be substantially unable to produce L-glutamic acid. Further, if *dtsR1* and *dtsR2* genes of coryneform L-glutamic acid producing bacteria having L-lysine producing ability are amplified, the bacteria are imparted with an ability to produce a marked amount of L-lysine (W095/23224, Japanese Patent Laid-open (Kokai) No. 10-234371/1998).

The *scrB* gene can be used for improvement of coryneform bacteria for use in the production of L-amino acids by using coryneform bacteria in a medium containing sucrose.

By deleting *aceA*, *accBC*, *lpd* or *odhA* of L-glutamic acid producing coryneform bacteria and so forth, their

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L-glutamic acid productivity can be enhanced. Further, *gluABCD* is a gene cluster of the L-glutamic acid uptake system, and by deleting one to four of *gluA*, *gluB*, *gluC* and *gluD* in coryneform L-glutamic acid producing

5 bacteria, the amount of L-glutamic acid accumulated in the medium can be increased. *aceA*, *accBC*, *lpd*, *odhA* and *gluABCD* of the present invention can be used for disruption of these genes on chromosome.

The medium used for producing L-amino acids by

10 utilizing a microorganism introduced with the DNA of the present invention may be a usual medium that contains a carbon source, a nitrogen source, inorganic ions and other organic trace nutrients as required. As the carbon source, there can be used hydrocarbons such as

15 glucose, lactose, galactose, fructose, sucrose, blackstrap molasses and starch hydrolysate; alcohols such as ethanol and inositol; or organic acids such as acetic acid, fumaric acid, citric acid and succinic acid.

As the nitrogen source, there can be used

20 inorganic ammonium salts such as ammonium sulfate, ammonium nitrate, ammonium chloride, ammonium phosphate and ammonium acetate, ammonia, organic nitrogen such as peptone, meat extract, yeast extract, corn steep liquor and soybean hydrolysate, ammonia gas, aqueous ammonia

25 and so forth.

As the inorganic ions (or sources thereof), added is a small amount of potassium phosphate, magnesium

sulfate, iron ions, manganese ions and so forth. As for the organic trace nutrients, it is desirable to add required substances such as vitamin B<sub>1</sub>, yeast extract and so forth in a suitable amount as required.

5           The culture is preferably performed under an aerobic condition attained by shaking, stirring for aeration or the like for 16 to 72 hours. The culture temperature is controlled to be at 30°C to 47°C, and pH is controlled to be 5 to 9 during the culture. As for  
10 the culture temperature, the culture may be performed at a temperature suitable for culture of a microorganism not introduced with the DNA of the present invention or a temperature higher than that. For adjustment of pH, inorganic or organic acidic or alkaline substances,  
15 ammonia gas and so forth can be used.

Collection of L-amino acids from fermentation broth can be attained by a combination of known methods such as techniques utilizing ion exchange resin, precipitation, crystallization and so forth depending on  
20 the kind of the L-amino acids.

#### Brief Explanation of the Drawings

Fig. 1 shows variation with temperature in activity of glutamate dehydrogenases derived from the  
25 *Corynebacterium thermoaminogenes* AJ12310 strain and the *Brevibacterium lactofermentum* 2256 strain.

Fig. 2 shows thermal stability of glutamate

dehydrogenases derived from the AJ12310 strain and the 2256 strain.

Fig. 3 shows variation with temperature in activity of citrate synthases derived from the AJ12310 strain and the 2256 strain.

Fig. 4 shows thermal stability of citrate synthases derived from the AJ12310 strain and the 2256 strain.

Fig. 5 shows variation with temperature in activity of isocitrate lyases derived from the AJ12310 strain and the 2256 strain.

Fig. 6 shows thermal stability of isocitrate lyases derived from the AJ12310 strain and the 2256 strain.

Fig. 7 shows variation with temperature in activity of phosphofructokinases derived from the AJ12310 strain and the 2256 strain.

Fig. 8 shows thermal stability of phosphofructokinases derived from the AJ12310 strain and the 2256 strain.

Fig. 9 shows variation with temperature in activity of phosphoenolpyruvate carboxylases derived from the AJ12310 strain and the 2256 strain.

Fig. 10 shows thermal stability of phosphoenolpyruvate carboxylases derived from the AJ12310 strain and the 2256 strain.

Fig. 11 shows variation with temperature in

activity of aconitases derived from the AJ12310 strain and the 2256 strain.

Fig. 12 shows thermal stability of aconitases derived from the AJ12310 strain and the 2256 strain.

5 Fig. 13 shows variation with temperature in activity of isocitrate dehydrogenases derived from the AJ12310 strain and the 2256 strain.

Fig. 14 shows thermal stability of isocitrate dehydrogenases derived from the AJ12310 strain and the  
10 2256 strain.

Fig. 15 shows thermal stability of 2-oxoglutarate dehydrogenases derived from the AJ12310 strain and the 2256 strain.

Fig. 16 shows construction of plasmid pSCR155  
15 carrying *scrB* gene.

Fig. 17 shows construction of plasmid pPDHA-2 carrying *pdhA* gene.

Fig. 18 shows L-glutamic acid productivity of a *pdhA* gene-amplified strain: (a) 37°C and (b) 44°C.

20 Fig. 19 shows is construction of a plasmid pICD-4 carrying *icd* gene.

Fig. 20 shows L-glutamic acid productivity of an *icd* gene-amplified strain: (a) 37°C and (b) 44°C.

Fig. 21 shows construction of plasmids pHSG299YGDH  
25 and pYGDH.

Fig. 22 shows construction of plasmids pHSG299YCS and pYCS.

Best Mode for Carrying out the Invention

Hereafter, the present invention will be further specifically explained with reference to the following  
5 examples.

Example 1

<1> Production of plasmid library of *Corynebacterium thermoaminogenes*

10           The *Corynebacterium thermoaminogenes* AJ12310 strain was cultured in CM2B liquid medium (1 g/dl of yeast extract (produced by Difco), 1 g/dl of polypeptone (produced by Nippon Seiyaku), 0.5 g/dl of NaCl, 10 µg/dl of biotin, pH 7.0 (adjusted with KOH)) at 37°C for 15  
15 hours, and its chromosomal DNA was prepared from the 10 ml of the medium by using a chromosomal DNA extraction kit (Bacterial Genome DNA Purification Kit (produced by Advanced Genetic Technologies)). The obtained DNA was partially digested with a restriction enzyme *Sau3AI*, and  
20 subjected to 0.8% agarose gel electrophoresis to fractionate the DNA. Then, a band corresponding to a DNA fragment of about 4 to 6 kb was excised from the gel, and a DNA fragment of the objective size was obtained by using a DNA gel extraction kit (GIBCO BRL, Concert™  
25 Rapid Gel Extraction System).

The plasmid pHSG399 (produced by Takara Shuzo) was fully digested with *BamHI*, and its end was



dephosphorylated by using alkaline phosphatase (CIAP; produced by Takara Shuzo). This vector fragment and the aforementioned chromosomal DNA fragment were ligated by using a DNA ligation kit produced by Takara Shuzo, and

5 *Escherichia coli* JM109 was the transformed with the obtained recombinant vector. Selection of transformants was performed on LB agar medium (containing 1.5 g/dl of agar) containing 30 µg/ml of chloramphenicol, 0.04 mg/ml of IPTG (isopropyl-β-D-thiogalactopyranoside) and 0.04

10 mg/ml of X-Gal (5-bromo-4-chloro-3-indolyl-β-D-galactoside) to obtain about 4000 white colonies.

<2> Design of primers for amplification of each gene

Primers for use in selection of a clone containing

15 each target gene by PCR from the plasmid library obtained above were designed. The target genes were mentioned above.

The primers were designed based on a known gene sequence of coryneform bacteria, i.e., its sequence of a

20 region where conservation at the amino acid level was observed when compared with corresponding genes of other microorganisms. Considering the codon usage of coryneform bacteria, a plurality of primer sets were designed for each gene.

25 To examine propriety of the prepared primers, PCR was performed by using these primers and chromosomal DNA of the *Corynebacterium thermoaminogenes* AJ12310 strain

as a template to amplify each gene fragment. As a result, when the PCR was performed by using the primers shown in the upper rows of Tables 2 to 7 under the conditions indicated as "PCR conditions for obtaining partial fragment" in the tables, an amplified fragment was observed for all of the genes. The parenthesized numbers after the primer sequences indicate the sequence numbers in Sequence Listing. These primers were used as primers for screening mentioned below.

Table 2

Gene	aceA	accBC	dtsR1
5'→3'Primer	CCTCTACCCAGCGAACTCCG (35)	CATCCACCCCGGTACGGCT (37)	ACGGCCAGCCCTGACCCGAC (39)
3'→5'Primer	CTGCCTTGAACCTACAGGTTC (36)	CGGTGACTGGGTGTTCCACC (38)	AGCAGCGCCCATGACGGCGA (40)
PCR conditions for obtaining partial fragment and PCR conditions for screening	94°C, 5 min 98°C, 5 sec 66°C, 2 sec, 30 cycles Z-Taq	94°C, 5 min 98°C, 5 sec 66°C, 2 sec, 30 cycles Z-Taq	94°C, 5 min 98°C, 5 sec 66°C, 2 sec, 30 cycles Z-Taq
Conditions of colony PCR	94°C, 7 min 91°C, 30 sec 55°C, 1 sec 72°C, 2.5 min, 30 cycles Ex-Taq	94°C, 7 min 91°C, 30 sec 55°C, 1 sec 72°C, 2.5 min, 30 cycles Ex-Taq	94°C, 7 min 91°C, 30 sec 55°C, 1 sec 72°C, 2.5 min, 30 cycles Ex-Taq
Amplified fragment	824bp	673bp	805bp

Table 3

Gene	dtsR2	pfk	scrB
5'→3'Primer	ACGGCCAGCCCTGACCGAC (41)	CGTCATCCGAGGAATCGTCC (43)	GGNCGHYTBAAYGAYCC (45)
3'→5'Primer	AGCAGCGCCCATGACGGCGA (42)	CGTGGCGGCCCATGACCTCC (44)	GGRCAYTCCCACATRTANCC (46)
PCR conditions for obtaining partial fragment and PCR conditions for screening	94°C, 5 min	94°C, 5 min	94°C, 5 min
	98°C, 5 sec	98°C, 5 sec	98°C, 5 sec
	66°C, 2 sec, 30 cycles	66°C, 2 sec, 30 cycles	50°C, 10 sec
	Z-Taq	Z-Taq	72°C, 20 sec, 40 cycles Z-Taq
Conditions of colony PCR	94°C, 7 min	94°C 7 min	94°C, 7 min
	91°C, 30 sec	91°C 30 sec	91°C, 30 sec
	55°C, 1 sec	55°C 1 sec	55°C, 1 sec
	72°C, 2.5 min, 30 cycles Ex-Taq	72°C 2.5 min 30 cycles Ex-Taq	72°C, 2.5 min, 30 cycles Ex-Taq
Amplified fragment	805bp	472bp	500bp

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Table 4

Gene	<i>gluABCD</i>	<i>pdhA</i>
5'→3'Primer	CCATCCGGATCCGGCAAGTC (47)	ACTGTGTCCATGGGTCTTGGCCC (49)
3'→5'Primer	AATCCCATCTCGTGGGTAAC (48)	CGCTGGAATCCGAACATCGA (50)
PCR conditions for obtaining partial fragment	94°C, 5 min 98°C, 5 sec 50°C, 10 sec 72°C, 20 sec, 30 cycles Z-Taq	94°C, 5 min 98°C, 5 sec 50°C, 10 sec 72°C, 20 sec, 30 cycles Z-Taq
Amplified fragment	500bp	1200bp
Conditions for screening PCR and colony PCR	94°C, 5 min 94°C, 30 sec 50°C, 1 min 72°C, 2 min, 30 cycles EX-Taq	94°C, 5 min 94°C, 30 sec 50°C, 1 min 72°C, 2 min, 30 cycles EX-Taq

Table 5

Gene	<i>pc</i>	<i>ppc</i>
5'→3'Primer	GGCGCAACCTACGACGTTGCAATGCG (51)	GGTTCCTGGATTGGTGGAGA(53)
3'→5'Primer	TGGCCGCCTGGGATCTCGTG (52)	CCGCCATCCTTGTTGGAATC(54)
PCR conditions for obtaining partial fragment	94°C, 5 min 98°C, 5 sec 55°C, 80 sec 30 cycles Z-Taq	94°C 5 min 98°C 5 sec 50°C 5 sec 72°C 10 sec 30 cycles Z-Taq
Amplified fragment	781bp	1000bp
Conditions for screening PCR	94°C, 5 min 98°C, 5 sec 55°C, 80 sec 30 cycles Z-Taq	94°C, 5 min 98°C, 5 sec 50°C, 5 sec 72°C, 10 sec, 30 cycles Z-Taq
Conditions for colony PCR	94°C, 5 min, 1 cycles 98°C, 5 sec 55°C, 80 sec, 50 cycles Z-Taq	94°C, 5 min 98°C, 5 sec 50°C, 10 sec 72°C, 20 sec, 50 cycles Z-Taq

Table 6

Gene	acn	icd	lpd
5'→3'Primer	GTIGGIACIGAYTCSCATAC (55)	GACATTTCACTCGCTGGACG (57)	ATCATCGCAACCGGTTTC (59)
3'→5'Primer	GCIGGAGAIATGTGRTCIGT (56)	CCGTACTCTTCAGCCTTCTG (58)	CGTCACCGATGGCGTAAAT (60)
PCR conditions	94°C, 1 min	94°C, 5 min	94°C, 5 min
for obtaining	96°C, 20 sec	98°C, 5 sec	98°C, 5 sec
partial	45°C, 1 min	55°C, 80 sec, 30 cycles	50°C, 10 sec
fragment	68°C, 2 min, 30 cycles	Z-Taq	72°C, 20 sec, 30 cycles
	EX-Taq		Z-Taq
Amplified fragment	1500bp	1500bp	500bp
Conditions for screening PCR and colony PCR	Same as above	Same as above	94°C, 5 min 94°C, 30 sec 57°C, 1 min 72°C, 1 min, 30 cycles Ex-Taq
Screening PCR			TACGAGGAGCAGATCCTCAA
5'→3'Primer			(63)
3'→5'Primer			TTGACGCCCGGTGTTCTCCAG
			(64)

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Table 6 (Cont.)

Gene	acn	icd	Lpd
LA cloning (N')	S1:GGTGAAGCTAAGTAGTACG 65) S2:AGCTACTAAACCTGCACC (66)	S1:CCGTACTCTTTCAGCCCTTCTG(67) S2:TCGTCTTGTTCACATC (68)	S1:ATCATCGCAACCGGTTT (69) S2:TACGAGGAGCAGATCCTCAA(70)
3'→5'Primer			
LA Cloning (C')	S1:GCTAACTACTTAGCTTCACC(71) S2:GAACCAGGAACCTATTGAACC(72)	S1:TCCGATGTCATCATCGAC (73) S2:ATGTGGAACAAGGACGAC (74)	
5'→3'Primer			
Restriction enzyme	PstI(N') HindIII(C')	Sali(N') PstI(C')	HindIII
Conditions for LA cloning	N' 94°C, 1 min 94°C, 30 sec 57°C, 2 min 72°C, 2 min, 30 cycles LA-Taq C' 94°C, 1 min 94°C, 30 sec 57°C, 2 min 72°C, 2.5 min, 30 cycles LA-Taq	94°C, 1 min 94°C, 30 sec 57°C, 2 min 72°C, 2.5 min, 30 cycles LA-Taq	94°C, 1 min 94°C, 30 sec 57°C, 2 min 72°C, 1 min, 30 cycles LA-Taq

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Table 7

Gene	<i>odhA</i>	
5'→3'Primer	ACACCGTGGTCGCCTCAACG (61)	
3'→5'Primer	TGCTAACCCGTCCCACCTGG (62)	
PCR conditions for obtaining partial fragment	94°C, 5 min 98°C, 5 sec 66°C, 2 sec, 30 cycles Z-Taq	
Amplified fragment	1306bp	
LA cloning (N') 5'→3'Primer	S1:GTACATATTGTCGTTAGAACGCGTAATACGACTCA(75) S2:CGTTAGAACGCGTAATACGACTCACTATAGGGAGA(76)	
Restriction	XbaI	
Conditions for LA cloning	<p>First time      94°C, 30 sec 55°C, 2 min 72°C, 1 min 30 cycles LA-Taq</p> <p>Second time      94°C, 1 min  98°C, 20 sec 68°C, 15 min, 30 cycles  72°C 10 min LA-Taq</p>	

### <3> Screening of plasmid library by PCR

A clone containing a target gene was selected from the plasmid library by PCR. Sixty colonies were picked up from each plasmid library, and replicated onto two LB agar medium plates. The 60 colonies of each plate were combined, inoculated to a test tube containing 4 ml of LB liquid medium and cultured for 15 hours. Then, a plasmid mixture was respectively obtained by using a plasmid DNA extraction kit produced by Promega. By using this plasmid mixture as a template and primers for screening prepared for each target gene, PCR was performed with the conditions shown as "conditions for screening PCR" in each table to select a clone from which a DNA fragment of the same size as that obtained by PCR using chromosomal DNA as a template had been amplified.

The nucleotide sequence of the amplified DNA fragment was determined by using a Big Dye dye terminator cycle sequencing kit produced by Perkin-Elmer, and investigating its homology to known gene information to determine if the target gene was obtained or not.

As for *lpd*, since any DNA fragment was not amplified with the primers produced in <2>, other primers for screening were prepared based on the determined nucleotide sequence.

### <4> Selection of clone harboring target gene by colony

## PCR

By using a plate that was an origin of a plasmid mixture for which amplification of the target gene fragment was confirmed, colony PCR was performed to select a clone containing the gene fragment. The colony PCR was performed with the conditions shown in Tables 2-7.

Plasmid DNA was collected from a selected transformant and the nucleotide sequence of the inserted DNA fragment was determined. When the full length of the target gene was not inserted in the inserted DNA fragment, and a upstream region, downstream region or the both were deleted, primers were prepared based on the determined nucleotide sequence, with which a gene fragment comprising the nucleotide sequence of the target gene in its full length was obtained by using TaKaRa LA PCR in vitro Cloning Kit (Takara Shuzo). Then, its nucleotide sequence was determined.

The outline of LA PCR cloning was as follows. Two kinds of primers each having one of the nucleotide sequences of two regions of the inserted DNA fragment were produced. Chromosomal DNA of *Corynebacterium thermoaminogenes* AJ12310 strain was digested with various restriction enzymes, and ligated to a cassette primer corresponding to each of the restriction enzymes. By using this as a template, PCR was performed with a primer (S1) corresponding to a position distant from the

deletion region and a cassette primer (C1) corresponding to a position outside the cassette primer among the prepared primers. Then, another PCR was performed with a primer (S2) corresponding to a position near the deletion region and a cassette primer (C2) corresponding to a position inside the cassette primer among the prepared primers. In this way, a DNA fragment containing the deleted region was obtained. By ligating the obtained DNA fragment with the already obtained DNA fragment, a DNA fragment containing the target gene in full length could be obtained. Since 5' end of the cassette did not have a phosphate group, a nick was formed at the ligation site of the 3' end of the DNA fragment and the 5' end of the cassette. Therefore, the DNA synthesis from the primer C1 stopped at this ligation site in the first PCR, and thus non-specific amplification did not occur. Therefore, specific amplification could be attained.

The primers and the reaction conditions used for the LA PCR cloning are shown in Tables 2-7. In the tables, the primers mentioned with "(N')" are primers used for the cloning of an upstream deleted portion, and the primers mentioned with "(C')" are primers used for the cloning of a downstream deleted portion. PCR was performed twice according to the instruction attached to the LA PCR cloning kit. Among the primers mentioned in the tables, the primers (S1) used for the first reaction

are shown in the upper row, and the primers (S2) used for the second reaction are shown in the lower row.

The nucleotide sequences of the DNA fragments containing each gene obtained as described above were  
5 determined in the same manner as mentioned above. Those nucleotide sequences and amino acid sequences that can be encoded by those nucleotide sequences are shown in SEQ ID NOS: 1-34. The sequences shown with the sequence numbers are summarized in Explanation of Sequence  
10 Listing mentioned hereinafter.

As for *scrB*, any open reading frame was not found. Since the *Corynebacterium thermoaminogenes* AJ12310 strain did not have the invertase activity and did not have sucrose assimilating property, an *scrB* gene  
15 fragment was obtained in a similar manner from *Corynebacterium thermoaminogenes* AJ12340 and AJ12309 strains having the sucrose assimilating property. As a result, a DNA fragment having an open reading frame was obtained from the both strains.

20

Example 2: Acquisition of *gdh* and *glta* gene

<1> Investigation of GDH activity of *Corynebacterium thermoaminogenes*

Cells of a wild strain of *Corynebacterium*  
25 *thermoaminogenes*, the AJ12310 strain, was grown on CM-2B agar medium (1 g/dl of yeast extract (produced by Difco), 1 g/dl of polypeptone (produced by Nippon Seiyaku), 0.5

g/dl of NaCl, 10  $\mu$ g/dl of biotin, 1.5 g/dl of agar, adjusted to pH 7.0 with KOH). The cells were inoculated to a 500-ml volume flask containing 20 ml of a medium for flask having the following composition and cultured at 37°C for 17 hours (until the residual sugar reached about 1 g/dl).

Similarly, cells of the 2256 strain (ATCC13869) of *Brevibacterium lactofermentum* grown on CM-2B agar medium were cultured at 31.5°C for 17 hours.

10

[Medium for flask]

Glucose	3 g/dl
KH <sub>2</sub> PO <sub>4</sub>	0.1 g/dl
MgSO <sub>4</sub> ·H <sub>2</sub> O	0.04 g/dl
15 FeSO <sub>4</sub> ·7H <sub>2</sub> O	1 mg/dl
MnSO <sub>4</sub> ·4H <sub>2</sub> O	1 mg/dl
Vitamin B <sub>1</sub> -HCl	200 $\mu$ g/L
Biotin	50 $\mu$ g/L
(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	1.5 g/dl
20 Soybean protein hydrolysis solution	48 mg/dl
(Memenno (T-N))	
CaCO <sub>3</sub> (Official reagent)	5 g/dl (separately sterilized)
pH 8.0 (adjusted with KOH)	

25

About 1 ml of the above culture medium was centrifuged at 1000 rpm for 1 minute to remove CaCO<sub>3</sub>, and the cells were washed twice with 200 mM K-phosphate

buffer (pH 6.9) and suspended in 300  $\mu$ l of the same buffer. The obtained cell suspension was sonicated for 5 minutes to disrupt the cells, centrifuged at 1000 rpm for 30 minutes to obtain a crude enzyme solution as the supernatant.

The optimum reaction temperature and the thermal stability of GDH activity were investigated using the aforementioned crude enzyme solution. The measurement of GDH activity was performed by adding the crude enzyme solution to a reaction mixture (100 mM Tris-HCl (pH 8.0), 20 mM  $\text{NH}_4\text{Cl}$ , 10 mM sodium  $\alpha$ -ketoglutarate, 0.25 mM NADPH) and measuring change of absorbance at 340 nm. The protein concentration of the crude enzyme solution was quantified by the Bradford method (Bio-Rad Protein Assay Kit was used) using bovine serum albumin as the standard through measurement of absorbance at 595 nm. The absorbance was measured by using HITACHI U-2000 (produced by Hitachi).

The GDH activity measured at various reaction temperatures is shown in Fig. 1. While the ATCC13869 strain showed the highest specific activity of GDH around 37°C and the activity markedly decreased around 42°C, the AJ12310 strain showed the highest specific activity around 42°C and it showed the activity even at 45°C.

Then, the thermal stability of GDH was investigated. The crude enzyme solution was left at

65°C for 0 to 30 minutes before the reaction, and then the enzyme activity was measured at 30°C. The results are shown in Fig. 2. As clearly seen from the results, while GDH of the ATCC13869 strain was inactivated by the heat treatment for 5 minutes, GDH of the AJ12310 strain maintained the activity even after the heat treatment for 30 minutes. In addition, the crude enzyme solution of the AJ12310 strain showed substantially no change in the GDH activity even after the heat treatment at 65°C for 90 minutes (data are not shown).

<2> Examination of CS activity of *Corynebacterium thermoaminogenes*

The optimum reaction temperature and thermal stability of CS were investigated by using crude enzyme solutions prepared from the cells of the *Corynebacterium thermoaminogenes* AJ12310 strain and the *Brevibacterium lactofermentum* ATCC13869 strain in the same manner as in Example 1. The measurement of CS activity was performed by adding each crude enzyme solution to a reaction mixture (100 mM Tris-HCl (pH 8.0), 0.1 mM DTNB (5,5'-dithiobis-(2-nitrobenzoic acid)), 200 mM sodium L-glutamate, 0.3 mM acetyl CoA), and measuring change of the absorbance at 412 nm.

The CS activity measured at various reaction temperatures is shown in Fig. 3. The ATCC13869 strain showed the highest specific activity of CS around 23°C



and the activity markedly decreased around 33°C. However, the AJ12310 strain showed high specific activity in a reaction temperature-dependent manner up to around 37°C and it showed the activity even at 40°C in a degree corresponding to about 40% of the activity at 37°C.

Then, thermal stability of CS was investigated. The crude enzyme solution was left at 33-55°C for 5 minutes before the reaction, and then the enzyme activity was measured at 30°C. The results are shown in Fig. 4. Whereas CS of the ATCC13869 strain was inactivated by the heat treatment at 35-40°C, CS of the AJ12310 strain maintained about 40% of the activity even after the heat treatment at 50°C.

### <3> Acquisition of *gdh* gene of *Corynebacterium thermoaminogenes*

The already reported nucleotide sequences of *gdh* gene of various microorganisms were compared. A region in which nucleotide sequences were well conserved was selected, and primers having the nucleotide sequences shown in SEQ ID NOS: 77 and 78 were prepared based on the nucleotide sequence of the region.

PCR was performed by using chromosomal DNA prepared from the *Corynebacterium thermoaminogenes* AJ12310 strain using Bacterial Genome DNA Purification Kit (produced by Advanced Genetic Technologies) as a

template and the aforementioned primers. Based on the obtained DNA fragment, genome walking was performed by using TaKaRa LA PCR in vitro Cloning Kit (produced by Takara Shuzo) to obtain the whole *gdh* gene, of which whole nucleotide sequence was determined. The result is shown in SEQ ID NO: 79. Further, the amino acid sequence deduced from this nucleotide sequence is shown in SEQ ID NO: 80.

The *gdh* gene of the *Brevibacterium lactofermentum* ATCC13869 strain was obtained in a similar manner, and its nucleotide sequence was determined. The result is shown in SEQ ID NO: 81. The amino acid sequence encoded by this nucleotide sequence is shown in SEQ ID NO: 82.

Homology was investigated for the nucleotide sequences of the *gdh* gene and the amino acid sequences of GDH of the *Corynebacterium thermoaminogenes* AJ12310 strain and the *Brevibacterium lactofermentum* ATCC13869 strain determined as described above, and the known *gdh* gene and amino acid sequence of GDH of the *Corynebacterium glutamicum* (*C. glutamicum*) ATCC13032 strain (Molecular Microbiology 6, 317-326 (1992)). The results are shown in Table 8 (for nucleotide sequences) and Table 9 (for amino acid sequences).

Table 8: Homology of nucleotide sequences of  
various *gdh* genes

	ATCC13869	ATCC13032	AJ12310
ATCC13869	-	94.5%	82.4%
ATCC13032	-	-	78.1%
AJ12310	-	-	-

Table 9: Homology of amino acid sequences of  
various GDH

	ATCC13869	ATCC13032	AJ12310
ATCC13869	-	90.8%	91.7%
ATCC13032	-	-	83.4%
AJ12310	-	-	-

#### <4> Acquisition of *gltA* gene of *Corynebacterium thermoaminogenes*

The already reported nucleotide sequences of *gltA* gene of various microorganisms were compared. A region in which nucleotide sequences were well conserved was selected, and primers having the nucleotide sequences shown in SEQ ID NOS: 83 and 84 were prepared based on the nucleotide sequence of the region.

PCR was performed by using chromosomal DNA prepared from the *Corynebacterium thermoaminogenes* AJ12310 strain (FERM BP-1542) using Bacterial Genome DNA Purification Kit (produced by Advanced Genetic Technologies) as a template and the aforementioned primers 7 and 8, and the nucleotide sequence of the amplified nucleotide sequence of about 0.9 kb was determined.

On the basis of the obtained nucleotide sequence of *gltA* gene of *Corynebacterium glutamicum* (Microbiol., 140, 1817-1828 (1994)), the primers of SEQ ID NOS: 85, 86, 87 and 88 were prepared. PCR was performed in a manner similar to the above by using chromosomal DNA of AJ12310 as a template and the primers of SEQ ID NOS: 85, 86, 87 and 88, and the nucleotide sequence of the amplified DNA fragment was specified to determine the whole nucleotide sequence of the *gltA* gene. The result is shown in SEQ ID NO: 89. Further, an amino acid sequence expected from this nucleotide sequence is shown in SEQ ID NO: 90.

The *gltA* gene of the *Brevibacterium lactofermentum* 2256 strain was obtained in a similar manner, and its nucleotide sequence was determined. The result is shown in SEQ ID NO: 91. The amino acid sequence encoded by this nucleotide sequence is shown in SEQ ID NO: 92.

Homology was investigated for the nucleotide sequences of the *gltA* gene and the amino acid sequences of CS of the *Corynebacterium thermoaminogenes* AJ12310 strain and the *Brevibacterium lactofermentum* ATCC13032 strain determined as described above, and the known *gltA* gene and amino acid sequence of CS of the *Corynebacterium glutamicum* ATCC13032 strain (Microbiol., 140, 1817-1828 (1994)). The results are shown in Table 10 (for nucleotide sequences) and Table 11 (for amino acid sequences).

Table 10: Homology of nucleotide sequences of  
various *gltA* genes

	ATCC13869	ATCC13032	AJ12310
ATCC13869	-	99.5%	85.7%
ATCC13032	-	-	85.6%
AJ12310	-	-	-

5 Table 11: Homology of amino acid sequences of  
various CS

	ATCC13869	ATCC13032	AJ12310
ATCC13869	-	99.3%	92.1%
ATCC13032	-	-	92.1%
AJ12310	-	-	-

Example 3: Acquisition of *scrB* gene of *Corynebacterium*  
*thermoaminogenes*

10 Since an *scrB* gene fragment was obtained from the  
*Corynebacterium thermoaminogenes* AJ12309 strain as shown  
in Example 1, it was attempted to obtain the total  
sequence of the gene. First, a partial fragment was  
obtained in the same manner as in Example 1 using the  
15 primers shown in SEQ ID NO: 45 and SEQ ID NO: 46. These  
primers were synthesized based on the *scrB* sequence of  
the *Brevibacterium lactofermentum* 2256 strain (Japanese  
Patent Laid-open No. 08-196280/1996).

20 Separately, chromosomal DNA was prepared from the  
AJ12309 strain by using Bacterial Genome DNA  
Purification Kit (Advanced Genetic Technologies Corp.).  
Sterilized water was added to 0.5  $\mu$ g of this chromosomal

DNA, 50 pmol each of the aforementioned primers, 4  $\mu$ l of dNTP mixture (2.5 mM each), 5  $\mu$ l of 10 x Z-Taq Buffer (Takara Shuzo) and 2 U of Z-Taq (Takara Shuzo) to prepare a PCR reaction mixture in a total volume of 50  $\mu$ l. PCR was performed with a cycle of denaturation at 98°C for 5 seconds, association at 50°C for 10 seconds and extension reaction at 72°C for 20 seconds, which was repeated for 30 cycles, by using the above reaction mixture and a thermal cycler GeneAmp PCR System 9600 (PE) to amplify a partial fragment of *scrB* of about 600 bp.

Then, the total sequence of *scrB* was determined by using an LA PCR in vitro Cloning Kit (Takara Shuzo). All of the procedure was performed in accordance with the protocol attached to the LA PCR in vitro Cloning Kit. Based on the obtained partial sequence, primers shown in SEQ ID NOS: 97, 98, 99 and 100 were synthesized. For the first PCR reaction for sequencing an upstream region, the primers shown in SEQ ID NOS: 95 and 97 and chromosomal DNA of AJ12309 strain digested with *Eco*T14I as a template DNA were used. For the second PCR reaction, the primers shown in SEQ ID NOS: 96 and 98 were used. For the first PCR reaction for sequencing a downstream region, the primers shown in SEQ ID NOS: 95 and 99 and chromosomal DNA of AJ12309 strain digested with *Sal*I (Takara Shuzo) as a template DNA were used. For the second PCR reaction, the primers shown in SEQ ID

NOS: 96 and 100 were used. By the above procedure, a sequence of a full length of 1656 bp containing ORF of *scrB* was determined. This nucleotide sequence is shown in SEQ ID NO: 93, and a deduced amino acid sequence is shown in SEQ ID NO: 94.

Example 4: Examination of thermal stability of isocitrate lyase, phosphofructokinase, phosphoenolpyruvate carboxylase, aconitase, isocitrate dehydrogenase and 2-oxoglutarate dehydrogenase

Thermal stability was investigated for the following enzymes derived from *Corynebacterium thermoaminogenes*. In this Example, protein concentrations were measured by the Bradford method (Bio-Rad Protein Assay Kit was used) using bovine serum albumin as a standard protein. Further, measurement of absorbance was performed by using HITACHI U-2000 (Hitachi) unless otherwise indicated.

<1> Isocitrate lyase

Thermal stability of activity of isocitrate lyase (henceforth also referred to as "ICL") derived from the *Corynebacterium thermoaminogenes* AJ12310 strain and ICL derived from the *Brevibacterium lactofermentum* 2256 strain (ATCC13869) was investigated. For the activity measurement, used were cells of which culture in a medium having the composition mentioned in Table 2 was

terminated before all of the carbon source was completely consumed. The method of the activity measurement was one described in Dieter J. Reinscheid et al., *J. Bacteriol.*, 176 (12), 3474 (1994). Specifically, the cells were washed with 50 mM Tris buffer (pH 7.3), suspended in the same buffer, and disrupted by sonication (INSONATOR 201M produced by KUBOTA was used, 200 W, 5 minutes). After the sonication, the suspension was centrifuged (13000 x g, 30 minutes) to remove undisrupted cells to prepare a crude enzyme solution.

The crude enzyme solution was added to a reaction system containing 50 mM MOPS-NaOH (pH 7.3), 5 mM dithiothreitol, 15 mM MgCl<sub>2</sub>, 1 mM EDTA, 5 mM D-threo-isocitrate, 0.2 mM NADH and 18 U of LDH (lactate dehydrogenase), and absorbance at 340 nm at various temperatures (30, 40, 50, 60 or 70°C) was measured by a Hitachi spectrophotometer U-3210. The measurement results for various reaction temperatures were shown in Fig. 5. Further, the crude enzyme solution was pretreated at 50°C (pretreatment time: 5 minutes or 15 minutes), and the activity was measured at 37°C. The results are shown in Fig. 6.

As a result, ICL of the AJ12310 strain showed the maximum activity at 60°C, whereas ICL of the 2256 strain showed the maximum activity around 50°C. Further, while ICL of the 2256 strain was completely inactivated after the pretreatment for 5 minutes, ICL of the AJ12310



strain maintained half of the activity after the pretreatment for 5 minutes. Thus, the stability of ICL of the AJ12310 strain at high temperatures was confirmed.

5                      Table 12   Composition of medium for  
ICL activity measurement

Component	Concentration
$(\text{NH}_4)_2\text{SO}_4$	5 g/l
Urea	5 g/l
$\text{KH}_2\text{PO}_4$	0.5 g/l
$\text{K}_2\text{HPO}_4$	0.5 g/l
MOPS	20.9 g/l
$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$	0.25 g/l
$\text{CaCl}_2 \cdot 7\text{H}_2\text{O}$	10 mM
$\text{CuSO}_4 \cdot 7\text{H}_2\text{O}$	0.2 mg/l
Biotin	0.2 mg/l
$\text{MnSO}_4 \cdot 7\text{H}_2\text{O}$	10 mg/l
$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$	10 mg/l
$\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$	1 mg/l
Acetic acid	4%

## <2> Phosphofructokinase

Thermal stability of activity of  
10    phosphofructokinase (henceforth also referred to as  
"PKF") derived from the *Corynebacterium thermoaminogenes*  
AJ12310 strain and PKF derived from the *Brevibacterium*  
*lactofermentum* 2256 strain was investigated. For the  
activity measurement, used were cells of which culture  
15    in a medium having the composition mentioned in Table 13  
was terminated before all of the saccharide was  
completely consumed. The method of the activity  
measurement was one described in Michiko Mori et al.,

*Agric. Biol. Chem.*, 51 (10), 2671 (1994). Specifically, the cells were washed with 0.1 M Tris buffer (pH 7.5), suspended in the same buffer, and disrupted by sonication (INSONATOR 201M produced by KUBOTA was used, 200 W, 5 minutes). After the sonication, the suspension was centrifuged (13000 x g, 30 minutes) to remove undisrupted cells to obtain a crude enzyme solution.

The crude enzyme solution was added to a reaction system containing 100 mM Tris buffer (pH 7.5), 0.2 mM NADH, 10 mM MgCl<sub>2</sub>, 2 mM NH<sub>4</sub>Cl, 10 mM KCl, 0.2 mM phosphoenolpyruvic acid, 6.4 mM fructose-6-phosphate, 1 mM ATP and 40 µg of LDH/PK (pyruvate kinase), and absorbance at 340 nm was measured at various temperatures (30, 40, 50, 60 or 70°C) by a Hitachi spectrophotometer U-3210. The measurement results for various reaction temperatures were shown in Fig. 7. Further, the crude enzyme solution was pretreated at 50°C (pretreatment time: 1, 3, 5 or 10 minutes), and the activity was measured at 37°C. The results are shown in Fig. 8.

As a result, PKF of the AJ12310 strain showed the maximum activity around 50°C, whereas PKF of the 2256 strain showed the maximum activity around 30°C. Thus, it was confirmed that the optimum temperature of PKF of the AJ12310 strain resided in a high temperature region.

Table 13 Composition of medium for  
PFK activity measurement

Component	Concentration
Polypeptone	20 g/l
Yeast extract	20 g/l
Sodium chloride	5 g/l
Glucose	20 g/l

### <3> Phosphoenolpyruvate carboxylase

5 Thermal stability of activity of  
phosphoenolpyruvate carboxylase (henceforth also  
referred to as "PEPC") derived from the *Corynebacterium*  
*thermoaminogenes* AJ12310 strain and PEPC of the  
*Brevibacterium lactofermentum* 2256 strain was examined.

10 Cells of the AJ12310 strain grown on CM-2B agar  
medium were inoculated to a 500-ml volume flask  
containing 20 ml of a medium for flask (8 g/dl of  
Glucose, 0.1 g/dl of  $\text{KH}_2\text{PO}_4$ , 0.04 g/dl of  $\text{MgSO}_4 \cdot \text{H}_2\text{O}$ , 1  
mg/dl of  $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ , 5 mg/dl of  $\text{MnSO}_4 \cdot 4\text{H}_2\text{O}$ , 3 g/dl of  
15  $(\text{NH}_4)_2\text{SO}_4$ , 48 mg/dl of TN (soybean protein hydrolysis  
solution), 200  $\mu\text{g/L}$  of vitamin  $\text{B}_1$ , 300  $\mu\text{g/L}$  of biotin,  
50  $\mu\text{l/l}$  of GD-113 (antifoaming agent), 5 g/dl of  $\text{CaCO}_3$   
(Official reagent, separately sterilized), pH 8.0  
(adjusted with KOH)), and cultured at 37°C. Cells of  
20 the 2256 strain grown on CM-2B agar medium were  
similarly cultured at 31.5°C.

The above culture broth in which the cells were  
grown to the logarithmic growth phase was centrifuged at  
1000 rpm for 1 minute to remove  $\text{CaCO}_3$ , and the cells

were washed 3 times with washing buffer (100 mM Tris/HCl pH 8.0, 10 mM  $\text{MgSO}_4$ , 1 mM DTT, 20% glycerol), sonicated to disrupt the cells, and centrifuged at 15 krpm for 10 minutes to remove cell debris. The supernatant was  
5 further centrifuged at 60 krpm for 1 hour to obtain a crude enzyme solution as the supernatant.

By using the above crude enzyme solution, optimum reaction temperature and thermal stability of the PEPC activity were investigated. The measurement of PEPC  
10 activity was performed by adding the crude enzyme solution to a reaction mixture (100 mM Tris/ $\text{H}_2\text{SO}_4$  (pH 8.5), 5 mM phosphoenolpyruvic acid, 10 mM  $\text{KHCO}_3$ , 0.1 mM acetyl-CoA, 0.15 mM NADH, 10 mM  $\text{MgSO}_4$ , 10 U of malate dehydrogenase, 0.1 mM DTT), and measuring change of the  
15 absorbance at 340 nm in 800  $\mu\text{l}$  of reaction volume.

The PEPC activity measured at various reaction temperatures is shown in Fig. 9. While the activity of the 2256 strain markedly decreased at 40°C, the AJ12310 strain showed substantially no decrease of the activity  
20 even at 40°C.

Then, the thermal stability of PEPC was investigated. The crude enzyme solution was left at 45°C for 0-20 minutes before the reaction, and then the enzyme activity was measured at 20°C. The results are  
25 shown in Fig. 10. As clearly seen from the results, whereas the PEPC activity of the 2256 strain was substantially lost after the heat treatment for 10

minutes, PEPC of the AJ12310 strain maintained the activity even after the heat treatment for 20 minutes.

These results demonstrated the stability of PEPC of the AJ12310 strain at a high temperature.

5

#### <4> Aconitase

Aconitase (henceforth also referred to as "ACN") derived from the *Corynebacterium thermoaminogenes* AJ12310 strain and ACN derived from the *Brevibacterium lactofermentum* 2256 strain were measured, and thermal stability thereof was examined.

Cells of the AJ12310 strain grown on CM-2B agar medium were inoculated to a 500-ml volume flask containing 20 ml of a medium for flask having the same composition as mentioned in <3>, and cultured at 37°C. Cells of the 2256 strain grown on CM-2B agar medium were similarly cultured at 31.5°C.

The above culture broth in which the cells were grown to the logarithmic growth phase was centrifuged at 1000 rpm for 1 minute to remove  $\text{CaCO}_3$ , and the cells were washed 3 times with 50 mM Tris/HCl pH 7.5, sonicated to disrupt the cells, and centrifuged at 15 krpm for 10 minutes to obtain a crude enzyme solution as the supernatant.

By using the above crude enzyme solution, optimum reaction temperature and thermal stability of ACN activity were investigated. The measurement of ACN

activity was performed by adding the crude enzyme solution to a reaction mixture (20 mM Tris/HCl (pH7.5), 50 mM NaCl, 20 mM isocitrate·3Na), and measuring change of the absorbance at 240 nm in 800 µl of reaction volume.

5           The ACN activity measured at various reaction temperatures is shown in Fig. 11. The AJ12310 strain showed higher activity at a higher temperature compared with the 2256 strain.

          Then, the thermal stability of ACN was  
10   investigated. The crude enzyme solution was left at 50°C for 0-15 minutes before the reaction, and then the enzyme activity was measured at 30°C. The results are shown in Fig. 12. As clearly seen from the results, ACN of the AJ12310 strain showed less activity decrease due  
15   to the heat treatment compared with ACN of the 2256 strain.

          These results demonstrated the stability of ACN of the AJ12310 strain at a high temperature.

#### 20   <5> Isocitrate dehydrogenase

          Thermal stability of activity of isocitrate dehydrogenase (henceforth also referred to as "ICDH") derived from the *Corynebacterium thermoaminogenes* AJ12310 strain and ICDH derived from the *Brevibacterium lactofermentum* 2256 strain was examined.  
25

          Cells of the AJ12310 strain grown on CM-2B agar medium were inoculated to a 500-ml volume flask

containing 20 ml of a medium for flask having the same composition as mentioned in <3>, and cultured at 37°C. Cells of the 2256 strain grown on CM-2B agar medium were similarly cultured at 31.5°C.

5           The above culture broth in which the cells were grown to the logarithmic growth phase was centrifuged at 1000 rpm for 1 minute to remove  $\text{CaCO}_3$ , and the cells were washed 3 times with 50 mM Tris/HCl pH 7.5, sonicated to disrupt the cells, and centrifuged at 15  
10       krpm for 10 minutes to obtain a crude enzyme solution as the supernatant.

          By using the above crude enzyme solution, optimum reaction temperature and thermal stability of ICDH activity were investigated. The measurement of ICDH  
15       activity was performed by adding the crude enzyme solution to a reaction mixture (35 mM Tris/HCl, 0.35 mM EDTA (pH 7.5), 1.5 mM  $\text{MnSO}_4$ , 0.1 mM NADP, 1.3 mM isocitrate·3Na), and measuring change of the absorbance at 340 nm in 800  $\mu\text{l}$  of reaction volume.

20           The ICDH activity measured at various reaction temperatures is shown in Fig. 13. While the activity of the 2256 strain markedly decreased at 70°C, substantially no activity decrease was observed even at 70°C for the AJ12310 strain.

25           Then, the thermal stability of ICDH was investigated. The crude enzyme solution was left at 45°C for 0-15 minutes before the reaction, and then the

enzyme activity was measured at 30°C. The results are shown in Fig. 14. As clearly seen from the results, while only about 15% of residual activity was observed after the heat treatment for 15 minutes for the 2256 strain, about 60% of residual ICDH activity was observed for the AJ12310 strain.

These results demonstrated the stability of ICDH of the AJ12310 strain at a high temperature.

#### 10 <6> 2-Oxoglutarate dehydrogenase

2-Oxoglutarate dehydrogenase (henceforth also referred to as "ODHC") derived from the *Corynebacterium thermoaminogenes* AJ12310 strain and ODHC derived from the *Brevibacterium lactofermentum* 2256 strain were measured, and thermal stability thereof was examined.

For the activity measurement, used were cells of which culture in a medium having the composition mentioned in Table 14 was terminated before all of the saccharide was completely consumed. The method of the activity measurement was one described in Isamu Shiio et al., Agric. Biol. Chem., 44 (8), 1897 (1980). Specifically, the cells were washed with 0.2% potassium chloride, suspended in 100 mM TES-NaOH (pH 7.5), 30% glycerol solution, and disrupted by sonication (INSONATOR 201M produced by KUBOTA was used, 200 W, 5 minutes). After the disruption by sonication, the suspension was centrifuged (13000 x g, 30 minutes) to



remove undisrupted cells, and subjected to gel filtration using the same buffer and Sephadex-G25 to prepare a crude enzyme solution.

The crude enzyme solution was added to a reaction system containing 100 mM TES-NaOH (pH 7.7), 5 mM  $\text{MgCl}_2$ , 0.2 mM Coenzyme A, 0.3 mM cocarboxylase, 1 mM  $\alpha$ -ketoglutaric acid, 3 mM L-cysteine and 1 mM acetylpyridine-adenine dinucleotide, and absorbance at 365 nm was measured at various temperatures (30, 40, 50, 60 or 70°C) by a Hitachi spectrophotometer U-3210. The crude enzyme solution was pretreated at 50°C (pretreatment time: 1, 3, 5 or 10 minutes), and the activity was measured at 37°C. The results are shown in Fig. 15.

As a result, while ODHC of the 2256 strain was completely inactivated by the pretreatment for 10 minutes, ODHC of the AJ12310 strain showed substantially constant activity irrespective of the pretreatment time, and thus its stability against high temperature treatment was confirmed.

Table 14 Composition of medium for  
ODHC activity measurement

Component	Concentration
Glucose	80 g/l
KH <sub>2</sub> PO <sub>4</sub>	1 g/l
MgSO <sub>4</sub> ·7H <sub>2</sub> O	0.4 g/l
FeSO <sub>4</sub> ·7H <sub>2</sub> O	0.01 g/l
MnSO <sub>4</sub> ·7H <sub>2</sub> O	0.05 g/l
(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	30 g/l
Soybean protein hydrolysate	480 mg/l
Thiamin hydrochloride	200 µg/l
Biotin	300 µg/l

Example 5: Impartation of sucrose assimilating ability  
5 by gene transfer of *scrB* gene

Since the *Corynebacterium thermoaminogenes* AJ12310 strain did not have invertase activity and sucrose assimilating property, it was investigated if sucrose assimilating ability could be imparted to it by  
10 transferring the *scrB* gene of the AJ12309 strain to the strain.

<1> Production of plasmid carrying *scrB* derived from  
*Corynebacterium thermoaminogenes* AJ12309 strain

15 To obtain an *scrB* gene fragment, the primers shown in SEQ ID NOS: 101 and 102 were synthesized, of which both ends were ligated with *Sma*I sequences, based on the nucleotide sequence shown in SEQ ID NO: 93. Sterilized water was added to 0.5 µg of chromosomal DNA of the  
20 12309 strain, 50 pmol each of the aforementioned

oligonucleotides, 4  $\mu$ l of dNTP mixture (2.5 mM each), 5  
5  $\mu$ l of 10 x Pyrobest Buffer (Takara Shuzo) and 2 U of  
Pyrobest polymerase (Takara Shuzo) to prepare a PCR  
reaction mixture in a total volume of 50  $\mu$ l. PCR was  
performed with a cycle of denaturation at 98°C for 10  
seconds, association at 55°C for 30 seconds and  
extension reaction at 72°C for 2 minutes, which was  
repeated for 30 cycles, by using the above reaction  
mixture and a thermal cycler GeneAmp PCR System 9600  
10 (PE) to amplify a fragment of about 1.7 kb containing  
*scrB* ORF.

Then, the above amplified fragment was digested  
with *Sma*I (Takara Shuzo), and ligated to plasmid pSAC4  
containing a dephosphorylated replication origin  
15 functioning in coryneform bacteria, which had been  
digested with *Sma*I, to prepare pSCR155. The  
construction of pSCR155 is shown in Fig. 16. pSAC4 was  
produced as follows. In order to make the vector for  
*Escherichia coli* pHSG399 (Takara Shuzo) autonomously  
20 replicable in coryneform bacteria, the replication  
origin (Japanese Patent Laid-open No. 5-7491/1993)  
derived from the already obtained plasmid pHM1519  
autonomously replicable in coryneform bacteria (Miwa,  
k.et al., Agric. Biol. Chem., 48 (1984) 2901-2903) was  
25 introduced into it. Specifically, pHM1519 was digested  
with restriction enzymes *Bam*HI and *Kpn*I, and the  
obtained fragment containing the replication origin was

blunt-ended by using a Blunting kit produced by Takara Shuzo and inserted into pHSG399 at the *SalI* site by using an *SalI* linker (produced by Takara Shuzo) to obtain pSAC4.

5

<2> Transfer of plasmid carrying *scrB* gene into AJ12310 strain

pSCR155 produced above and plasmid pSSM30BS (Japanese Patent Laid-open No. 08-196280/1996) carrying the *scrB* gene derived from *Brevibacterium lactofermentum* were introduced into the *Corynebacterium thermoaminogenes* AJ12310 strain. The transformation was performed according to the following procedure. The cells were inoculated to CM-2B medium containing 20% sucrose in such an amount that OD<sub>660</sub> of the medium should become 0.1, and cultured at 37°C with shaking until the OD<sub>660</sub> become 0.3. Lysozyme was added to the medium at a concentration of 100 µg/ml, and the cells were further cultured for 2 hours. The cells were washed three times with 20% sucrose, suspended in 20% sucrose, added with the plasmid collected from *Escherichia coli* JM110, mixed sufficiently, and applied with an electric pulse (18 kV/cm, 300 msec) to be introduced with the DNA. After the cells were subjected to restoration culture overnight in CM-2B medium containing 20% sucrose, transformants were selected on CM-2B agar medium containing 5 µg/ml of chloramphenicol. Specifically,

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20

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the transformation was performed by the electric pulse method (Japanese Patent Laid-open No. 12-204236/2000, and the selection of transformants was performed on CM2B plate medium containing 5 µg/ml of chloramphenicol at  
5 37°C. As a result, any transformant harboring the plasmid pSSM30BS carrying *scrB* derived from *Brevibacterium lactofermentum* was not obtained, but only a transformant harboring the plasmid pSCR155 carrying *scrB* derived from *Corynebacterium thermoaminogenes* was  
10 obtained. This strain was designated as AJ12310/pSCR155.

<3> Evaluation of culture of AJ12310/pSCR155 strain using sucrose as sugar source.

AJ12310/pSCR155 prepared above was inoculated to a  
15 medium having the composition shown in Table 15, and cultured at 37°C for 22 hours with shaking. The absorbance (OD) and residual sugar (RS) of the medium were measured after the culture. The results are shown in Table 16. As a result, it was confirmed that, while  
20 the AJ12310 strain could not assimilate sucrose and hence could not grow, the *scrB* gene introduced strain, the AJ12310/pSCR155 strain, became to be able to assimilate sucrose.

Table 15 Medium composition

Medium composition	Concentration
Sucrose	60 g/l
$\text{KH}_2\text{PO}_4$	1 g/l
$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$	0.4 g/l
$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
$\text{MnSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
$(\text{NH}_4)_2\text{SO}_4$	30 g/l
Soybean protein hydrolysate	480 mg/l
Thiamin hydrochloride	200 $\mu\text{g/l}$
Biotin	300 $\mu\text{g/l}$

Table 16 Result of sucrose culture

	OD (x 51)	RS (g/l)
2256	1.292	0.00
AJ12310	0.058	60.00
AJ12310/pSCR155	1.571	0.84

5     Example 6: L-glutamic acid production by *pdhA* gene-amplified strain

<1> Construction of plasmid pPDHA-2 carrying *pdhA*

10     The *pdhA* gene derived from the *Corynebacterium thermoaminogenes* AJ12310 strain was obtained by screening of a plasmid library. Specifically, PCR was performed with the conditions shown in Example 1, Table 4, using a plasmid library mixture as a template, and a clone p21A was selected, from which a DNA fragment of the same size is amplified as obtained in PCR using

15     chromosomal DNA as a template. The DNA sequence of this plasmid was determined to confirm that the full length of *pdhA* was contained in it.

p21A was digested with *Xba*I and *Kpn*I to excise a DNA fragment of 4 kb containing the full length of the *pdhA* gene and a promoter region. This DNA fragment containing the *pdhA* gene was inserted into the *Xba*I and *Kpn*I sites of pHSG299 (Takara Shuzo). Then, this plasmid was digested with *Xba*I, and a fragment obtained by digesting pXK4 with *Xba*I was inserted to prepare pPDHA-2. The construction process of pPDHA-2 is shown in Fig. 17. A DNA Ligation Kit Ver.2 (Takara Shuzo) was used for the ligation reaction, and *Escherichia coli* JM109 strain (Takara Shuzo) was used as the host of genetic manipulation. The aforementioned pXK4 was produced as follows. A shuttle vector pHK4 for coryneform bacteria and *Escherichia coli* (Japanese Patent Laid-open No. 5-7491/1993) was digested with restriction enzymes *Bam*HI and *Kpn*I to obtain a DNA fragment containing the replication origin, and the obtained fragment was blunt-ended by using a DNA blunting kit (Blunting Kit produced by Takara Shuzo), ligated to an *Xba*I linker (produced by Takara Shuzo) and inserted into pHSG299 at the *Xba*I site to obtain the plasmid pXK4.

<2> Transfer of plasmid carrying *pdhA* gene into AJ12310 strain

The plasmid pPDHA-2 produced above was introduced into the *Corynebacterium thermoaminogenes* AJ12310 strain

to prepare a *pdhA* gene-amplified strain. The transformation was performed in the same manner as Example 5, and a transformant was selected on CM-2B agar medium containing 25  $\mu$ g/ml kanamycin to obtain

5 AJ12310/pPDHA-2 strain.

<3> L-glutamic acid production by *pdhA*-amplified strain

The AJ12310 strain and the *pdhA* gene-amplified strain obtained above, AJ12310/pPDHA-2 strain, both of which were grown on CM-2B agar medium, were each inoculated to a 500-ml volume flask containing 20 ml of a medium for seed culture flask shown in Table 17, and cultured at 37°C with shaking until glucose was completely consumed. 2 ml of this culture broth was

10 inoculated into 500 ml-volume flask containing 20 ml of a medium for main culture flask shown in Table 17, and cultured as main culture at 37°C and 44°C. The main culture was continued until glucose was completely consumed. After the culture, OD<sub>620</sub> of the medium and

15 accumulated amount of L-glutamic acid were measured to examine the effect of the gene amplification on the cell formation and production of glutamic acid. The measurement of OD was performed by using a spectrophotometer HITACHI U-2000 (Hitachi), and L-

25 glutamic acid concentration was measured by using a glutamic acid analyzer AS-210 (Asahi Chemical Industry). The results are shown in Fig. 18.



The *pdhA* gene-amplified strain, AJ12310/pPDHA-2 strain, showed increased L-glutamic acid accumulation and increased OD compared with the AJ12310 strain, and thus it became clear that the amplification of the *pdhA* gene was effective for L-glutamic acid production.

Table 17 Medium for evaluation of  
*pdhA*-amplified strain

Medium composition	Seed culture	Main culture
Sucrose	30 g/l	60 g/l
KH <sub>2</sub> PO <sub>4</sub>	1 g/l	1 g/l
MgSO <sub>4</sub> ·7H <sub>2</sub> O	0.4 g/l	0.4 g/l
FeSO <sub>4</sub> ·7H <sub>2</sub> O	0.01 g/l	0.01 g/l
MnSO <sub>4</sub> ·7H <sub>2</sub> O	0.01 g/l	0.01 g/l
(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	15 g/l	30 g/l
Soybean protein hydrolysate	480 mg/l	480 mg/l
Thiamin hydrochloride	200 µg/l	200 µg/l
Biotin	10 µg/l	
AZ-20R (anti-foaming agent)	20 µg/l	20 µg/l
CaCO <sub>3</sub> (separately sterilized)	50 g/L	50 g/L
pH 8.0 (adjusted with KOH)		

#### 10 Example 7: L-glutamic acid production by *icd* gene-amplified strain

<1> Construction of plasmid pICD-4 carrying *icd* derived from *Corynebacterium thermoaminogenes* AJ12310 strain

Based on the *icd* gene sequence of the AJ12310 strain shown in SEQ ID NO: 29, the primers shown in SEQ ID NO: 103 and SEQ ID NO: 104 were synthesized. A *Bgl*III site was introduced into 5' end of the both primers.

Separately, genomic DNA of the *Corynebacterium thermoaminogenes* AJ12310 strain was prepared by using a Genomic DNA Purif. Kit (Edge BioSystems). Sterilized water was added to the genome DNA as a template, 100  
5 pmol each of the aforementioned primers, 8  $\mu$ l of dNTP mixture (2.5 mM each), 10  $\mu$ l of 10 x Pyrobest Buffer II (Takara Shuzo) and 2.5 U of Pyrobest polymerase (Takara Shuzo) to prepare a PCR reaction mixture in a total volume of 100  $\mu$ l. PCR was performed with a cycle of  
10 denaturation at 98°C for 10 seconds, association at 55°C for 1 minute and extension reaction at 72°C for 4 minutes, which was repeated for 30 cycles, by using the above reaction mixture and a thermal cycler TP240 (Takara Shuzo) to amplify a DNA fragment of about 3.3 kb  
15 containing the *icd* gene and its promoter.

Then, this DNA fragment containing the *icd* gene was digested with *Bgl*III, and ligated to pHSG299 (Takara Shuzo) at the *Bam*HI site. This plasmid was then treated with *Xba*I, and a fragment obtained by digesting pXK4  
20 with *Xba*I was inserted into the plasmid to construct pICD-4. The construction procedure of pICD-4 is shown in Fig. 19. A DNA Ligation Kit Ver.2 (Takara Shuzo) was used for the ligation reaction, and *Escherichia coli* JM109 strain (Takara Shuzo) was used as the host of  
25 genetic manipulation.

<2> Transfer of plasmid carrying *icd* gene into AJ12310

strain

The plasmid pICD-4 produced above was introduced into the *Corynebacterium thermoaminogenes* AJ12310 strain to prepare an *icd* gene-amplified strain. The transformation was performed in the same manner as Example 5, and a transformant was selected on CM-2B agar medium containing 25 µg/ml kanamycin to obtain AJ12310/pICD-4 strain.

<3> L-glutamic acid production by *icd*-amplified strain

Culture evaluation was performed for the AJ12310 strain and the *icd*-amplified strain thereof, AJ12310/pICD, by the culture method described in Example 6. The results are shown in Fig. 20. The *icd* gene-amplified strain AJ12310/pICD-4 strain showed increased L-glutamic acid accumulation and increased OD compared with the AJ12310 strain, and thus it became clear that the amplification of the *icd* gene was effective for L-glutamic acid production.

Example 8: L-glutamic acid production by *gdh* gene-amplified strain

<1> Construction of plasmid carrying *gdh* derived from *Corynebacterium thermoaminogenes* AJ12310 strain

Based on the *gdh* gene sequence of the AJ12310 strain shown in SEQ ID NO: 79, the primers shown in SEQ ID NO: 105 and SEQ ID NO: 106 were synthesized.

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arately, chromosomal DNA of the AJ12310 strain was prepared by using a Bacterial Genome DNA Purification Kit (Advanced Genetic Technologies Corp.). Sterilized water was added to 0.5  $\mu$ g of this chromosomal DNA, 10 pmol each of the aforementioned oligonucleotides, 8  $\mu$ l of dNTP mixture (2.5 mM each), 5  $\mu$ l of 10 x LA Taq Buffer (Takara Shuzo) and 2 U of LA Taq (Takara Shuzo) to prepare a PCR reaction mixture in a total volume of 50  $\mu$ l. PCR was performed with a cycle of denaturation at 94°C for 30 seconds, association at 55°C for 1 second and extension reaction at 72°C for 3 minutes, which was repeated for 30 cycles, by using the above reaction mixture and a thermal cycler TP240 (Takara Shuzo) to amplify a DNA fragment of about 2 kb containing the *gdh* gene and its promoter. The obtained amplified fragment was digested with *Pst*I (Takara Shuzo), mixed with pHSG299 (Takara Shuzo) fully digested with *Pst*I and ligated to it. A DNA Ligation Kit Ver.2 produced by Takara Shuzo was used for the ligation reaction. After the ligation, competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo) were transformed with the ligation product, plated on L medium (10 g/l of Bacto-trypton, 5 g/l of Bacto-yeast extract, 5 g/l of NaCl, 15 g/l of agar, pH 7.2) containing 10  $\mu$ g/ml of IPTG (isopropyl- $\beta$ -D-thiogalactopyranoside), 40  $\mu$ g/ml of X-Gal (5-bromo-4-chloro-3-indolyl- $\beta$ -D-galactoside) and 40

$\mu\text{g/ml}$  of chloramphenicol, and cultured overnight. The emerged white colonies were picked up and subjected to single colony separation to obtain transformants.

Plasmids were prepared from the transformants by the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992) and their restriction maps were prepared. A plasmid having a restriction map equivalent to that shown in Fig. 21 was designated as pHSG299YGDH.

A replication origin that functions in coryneform bacteria was introduced into this pHSG299YGDH. Specifically, pXC4 was digested with a restriction enzyme *Xba*I to obtain a fragment containing a replication origin derived from pHM1519, and it was mixed with pHSG299YGDH fully digested with *Xba*I and ligated to it. Plasmids were prepared in the same manner as above and a plasmid having a restriction map equivalent to that shown in Fig. 21 was designated as pYGDH. pXC4 was constructed in the same manner as that for pXK4 mentioned in Example 6 except that pHSG399 ( $\text{Cm}^r$ ) was used instead of pHSG299.

<2> Transfer of plasmid carrying *gdh* gene into AJ12310

The plasmid produced above was introduced into the *Corynebacterium thermoaminogenes* AJ12310 strain to prepare a *gdh* gene-amplified strain. The transformation

was performed in the same manner as Example 5, and a transformant was selected on CM-2B agar medium containing 25  $\mu$ g/ml kanamycin at 31°C to obtain AJ12310/pYGDH.

5

<3> L-glutamic acid production by *gdh*-amplified strain

The AJ12310 strain and the *gdh* gene-amplified strain obtained above, AJ12310/pYGDH strain, both of which were grown on CM-2B agar medium, were each  
10 inoculated to a 500-ml volume flask containing 20 ml of a medium for seed culture flask shown in Table 18, and cultured at 37°C with shaking until glucose was completely consumed. 2 ml of this culture broth was inoculated into 500 ml-volume flask containing 20 ml of  
15 a medium for main culture flask shown in Table 19, and cultured as main culture at 37°C and 44°C. The main culture was continued until glucose was completely consumed. After completion of the culture,  $OD_{620}$  of the medium and accumulated amount of L-glutamic acid were  
20 measured to examine the effect of the gene amplification on the cell formation and production of glutamic acid. The measurement of OD was performed by using a spectrophotometer HITACHI U-2000 (Hitachi), and L-glutamic acid concentration was measured by using a  
25 glutamic acid analyzer AS-210 (Asahi Chemical Industry).

Table 18 Composition of medium  
for seed culture

Medium composition	Concentration
Glucose	30 g/l
Ammonium sulfate	15 g/l
$\text{KH}_2\text{PO}_4$	1 g/l
$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$	0.4 g/l
$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
$\text{MnSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
Soybean protein hydrolysate	0.48 g/l
Thiamin hydrochloride	200 $\mu\text{g/l}$
Biotin	10 $\mu\text{g/l}$
AZ20R	0.02 ml/l
$\text{CaCO}_3$ (separately sterilized)	1 g/L
pH 8.0 (adjusted with KOH)	

Table 19 Composition of medium  
for main culture

Medium composition	Concentration
Glucose	60 g/l
Ammonium sulfate	30 g/l
$\text{KH}_2\text{PO}_4$	1 g/l
$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$	0.4 g/l
$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
$\text{MnSO}_4 \cdot 7\text{H}_2\text{O}$	0.01 g/l
Soybean protein hydrolysate	0.48 g/l
Thiamin hydrochloride	200 $\mu\text{g/l}$
AZ20R	0.02 ml/l
$\text{CaCO}_3$ (separately sterilized)	1 g/L
PH 8.0 (adjusted with KOH)	

The results of the culture are shown in Table 20 and Table 21. At 37°C, the *gdh*-amplified strain showed higher saccharide consuming rate, better growth and

higher attained OD compared with the parent strain, the AJ12310 strain. Moreover, both of the L-glutamic acid accumulation and the yield were markedly improved, i.e., 5-7%, at 37°C. Also at 44°C, the yield was improved, and the attained OD increased. On the other hand, it was confirmed that accumulation of  $\alpha$ -ketoglutaric acid was decreased in the *gdh*-amplified strain. These results demonstrate that the amplification of *gdh* is effective for improvement in L-glutamic acid yield and reduction of byproduct.

Table 20 Culture result of *gdh*-amplified strain (37°C)

	OD <sub>620</sub> (51x)	L-Glu accumulation (g/dl)	L-Glu yield (%)	$\alpha$ -KG (mg/dl)
AJ12310	0.58	1.74	30.7	53.9
AJ12310/PYGDH	0.65	2.23	39.3	4.1

Table 21 Culture result of *gdh*-amplified strain (44°C)

	OD <sub>620</sub> (51x)	L-Glu accumulation (g/dl)	L-Glu yield (%)
AJ12310	0.63	1.70	26.7
AJ12310/pYGDH	0.71	1.79	27.8

#### Example 9: L-glutamic acid production by *gltA* gene-amplified strain

<1> Construction of plasmid carrying *gltA* gene derived from *Corynebacterium thermoaminogenes*

Based on the *gltA* gene sequence of the AJ12310 strain shown in SEQ ID NO: 89, the primers shown in SEQ



ID NO: 107 and SEQ ID NO: 108 were synthesized.

Separately, chromosomal DNA of the AJ12310 strain was prepared by using a Bacterial Genome DNA Purification Kit (Advanced Genetic Technologies Corp.).

5 Sterilized water was added to 0.5  $\mu$ g of this chromosomal DNA, 10 pmol each of the aforementioned oligonucleotides, 8  $\mu$ l of dNTP mixture (2.5 mM each), 10  $\mu$ l of 10 x Pyrobest-Taq Buffer (Takara Shuzo) and 2 U of Pyrobest Taq (Takara Shuzo) to prepare a PCR reaction mixture in

10 a total volume of 100  $\mu$ l. PCR was performed with a cycle of denaturation at 94°C for 30 seconds, association at 45°C for 30 seconds and extension reaction at 72°C for 3 minutes, which was repeated for 30 cycles, by using the above reaction mixture and a

15 thermal cycler TP240 (Takara Shuzo) to amplify a DNA fragment of about 2 kb containing the *gltA* gene and its promoter. The obtained amplified fragment was digested with *Kpn*I (Takara Shuzo), mixed with pHSG299 (Takara Shuzo) fully digested with *Kpn*I and ligated to it. A

20 DNA Ligation Kit Ver.2 produced by Takara Shuzo was used for the ligation reaction. After the ligation, competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo) were transformed with the ligation product, plated on L medium (10 g/l of Bacto-trypton, 5 g/l of

25 Bacto-yeast extract, 5 g/l of NaCl, 15 g/l of agar, pH 7.2) containing 10  $\mu$ g/ml of IPTG (isopropyl- $\beta$ -D-thiogalactopyranoside), 40  $\mu$ g/ml of X-Gal (5-bromo-4-

chloro-3-indolyl- $\beta$ -D-galactoside) and 40  $\mu$ g/ml of chloramphenicol, and cultured overnight. The emerged white colonies were picked up and subjected to single colony separation to obtain transformants.

5           Plasmids were prepared from the transformants by the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992) and their restriction maps were prepared. A plasmid having a restriction map  
10           equivalent to that shown in Fig. 22 was designated as pHSG299YCS.

          A replication origin that is replicable in coryneform bacteria was introduced into this pHSG299YCS. Specifically, pXC4 was digested with a restriction  
15           enzyme *Xba*I to obtain a fragment containing a replication origin derived from pHM1519, and it was mixed with pHSG299YCS fully digested with *Xba*I and ligated to it. Plasmids were prepared in the same  
20           manner as above and a plasmid having a restriction map equivalent to that shown in Fig. 22 was designated as pYCS.

<2> Transfer of plasmid carrying *gltA* gene into AJ12310 strain

25           The plasmid produced above was introduced into the *Corynebacterium thermoaminogenes* AJ12310 strain to prepare a *gltA* gene-amplified strain. The

transformation was performed in the same manner as Example 5, and a transformant was selected on CM-2B agar medium containing 25  $\mu$ g/ml kanamycin to obtain AJ12310/pYCS.

5

<3> L-glutamic acid production by *gltA*-amplified strain

The AJ12310 strain and the *gltA* gene-amplified strain obtained above, AJ12310/pYCS strain, both of which were grown on CM-2B agar medium, were cultured in the same manner as in Example 8. The results of the culture are shown in Table 22 and Table 23. Both at the culture temperatures, 37°C and 44°C, the CS-enhanced strain showed improved glutamic acid accumulation compared with the parent strain. Further, the *gltA*-amplified strain showed decreased L-aspartic acid and L-lysine, which are synthesized from oxaloacetic acid.

10  
15

These results demonstrate that the amplification of *gltA* is effective for improvement of L-glutamic acid yield and reduction of byproduct.

20

Table 22 Culture result of *gltA*-amplified strain (37°C)

	L-Glu accumulation (g/dl)	Yield (%)	L-Asp accumulation (mg/dl)	L-Lys accumulation (mg/dl)
AJ12310	1.79	31.9	11.8	11.0
AJ12310/pYCS	2.04	36.5	8.1	7.3

Table 23 Culture result of *gluA*-amplified strain (44°C)

	OD	L-Glu accumulation (g/dl)	Yield (%)	L-Asp accumulation (mg/dl)	L-Lys Accumulation (mg/dl)
AJ12310	0.58	1.38	21.8	23.3	29.2
AJ12310/pYCS	0.65	1.84	28.8	14.1	17.2

## [Explanation of Sequence Listing]

- SEQ ID NO: 1: *aceA*, nucleotide sequence
- SEQ ID NO: 2: *aceA*, amino acid sequence
- SEQ ID NO: 3: *accBC*, nucleotide sequence
- SEQ ID NO: 4: *accBC*, amino acid sequence
- SEQ ID NO: 5: *dtsR1*, nucleotide sequence
- SEQ ID NO: 6: *dtsR1*, amino acid sequence
- SEQ ID NO: 7: *dtsR2*, nucleotide sequence
- SEQ ID NO: 8: *dtsR2*, amino acid sequence
- SEQ ID NO: 9: *pfk*, nucleotide sequence
- SEQ ID NO: 10: *pfk*, amino acid sequence
- SEQ ID NO: 11: *scrB* (AJ12340), nucleotide sequence
- SEQ ID NO: 12: *scrB* (AJ12340), amino acid sequence
- SEQ ID NO: 13: *scrB* (AJ12309), nucleotide sequence
- SEQ ID NO: 14: *scrB* (AJ12309), amino acid sequence
- SEQ ID NO: 15: *scrB* (AJ12310), nucleotide sequence
- SEQ ID NO: 16: *gluABCD*, nucleotide sequence
- SEQ ID NO: 17: *gluABCD*, amino acid sequence
- SEQ ID NO: 18: *gluABCD*, amino acid sequence
- SEQ ID NO: 19: *gluABCD*, amino acid sequence
- SEQ ID NO: 20: *gluABCD*, amino acid sequence
- SEQ ID NO: 21: *pdhA*, nucleotide sequence

SEQ ID NO: 22: *pdhA*, amino acid sequence  
SEQ ID NO: 23: *pc*, nucleotide sequence  
SEQ ID NO: 24: *pc*, amino acid sequence  
SEQ ID NO: 25: *ppc*, nucleotide sequence  
SEQ ID NO: 26: *ppc*, amino acid sequence  
SEQ ID NO: 27: *acn*, nucleotide sequence  
SEQ ID NO: 28: *acn*, amino acid sequence  
SEQ ID NO: 29: *icd*, nucleotide sequence  
SEQ ID NO: 30: *icd*, amino acid sequence  
SEQ ID NO: 31: *lpd*, nucleotide sequence  
SEQ ID NO: 32: *lpd*, amino acid sequence  
SEQ ID NO: 33: *odhA*, nucleotide sequence  
SEQ ID NO: 34: *odhA*, amino acid sequence  
SEQ ID NO: 79: *gdh* (AJ12310), nucleotide sequence  
SEQ ID NO: 80: *gdh* (AJ12310), amino acid sequence  
SEQ ID NO: 81: *gdh* (2256), nucleotide sequence  
SEQ ID NO: 82: *gdh* (2256), amino acid sequence  
SEQ ID NO: 89: *glta* (AJ12310), nucleotide sequence  
SEQ ID NO: 90: *glta* (AJ12310), amino acid sequence  
SEQ ID NO: 91: *glta* (2256), nucleotide sequence  
SEQ ID NO: 92: *glta* (2256), amino acid sequence  
SEQ ID NO: 93: *scrB* (AJ12309), nucleotide sequence  
SEQ ID NO: 94: *scrB* (AJ12309), amino acid sequence

#### Industrial Applicability

According to the present invention, genes coding

for enzymes of amino acid biosynthetic pathway derived from *Corynebacterium thermoaminogenes*, or genes coding for proteins involved in the amino acid uptake into cells.

The genes of the present invention can be utilized for the production of the aforementioned enzymes or proteins, or the breeding of amino acid producing bacteria.

What is claimed is:

1. A protein having the amino acid sequence of SEQ ID NO: 2 or the amino acid sequence of SEQ ID NO: 2 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has isocitrate lyase activity and shows 30% or more of residual activity after a heat treatment at 50°C for 5 minutes.

2. A protein having the amino acid sequence of SEQ ID NO: 4 or the amino acid sequence of SEQ ID NO: 4 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which is involved in acyl Co-A carboxylase activity and is derived from *Corynebacterium thermoaminogenes*.

3. A protein having the amino acid sequence of SEQ ID NO: 6 or the amino acid sequence of SEQ ID NO: 6 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has DtsR activity and is derived from *Corynebacterium thermoaminogenes*.

4. A protein having the amino acid sequence of SEQ ID NO: 8 or the amino acid sequence of SEQ ID NO: 8

including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has DtsR activity and is derived from *Corynebacterium thermoaminogenes*.

5. A protein having the amino acid sequence of SEQ ID NO: 10 or the amino acid sequence of SEQ ID NO: 10 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which shows phosphofructokinase activity at 60°C in an equivalent or higher degree compared with the activity at 30°C.

6. A protein having the amino acid sequence of SEQ ID NO: 94 or the amino acid sequence of SEQ ID NO: 94 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has activity for imparting sucrose assimilating ability to *Corynebacterium thermoaminogenes*.

7. A protein having any one of the amino acid sequences of SEQ ID NOS: 17-20 or the amino acid sequence of any one of SEQ ID NOS: 17-20 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has a function involved in glutamic acid uptake and is derived from *Corynebacterium thermoaminogenes*.



8. A protein having the amino acid sequence of SEQ ID NO: 22 or the amino acid sequence of SEQ ID NO: 22 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has pyruvate dehydrogenase activity and is derived from *Corynebacterium thermoaminogenes*.

9. A protein having the amino acid sequence of SEQ ID NO: 24 or the amino acid sequence of SEQ ID NO: 24 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has pyruvate carboxylase activity and is derived from *Corynebacterium thermoaminogenes*.

10. A protein having the amino acid sequence of SEQ ID NO: 26 or the amino acid sequence of SEQ ID NO: 26 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has phosphoenolpyruvate carboxylase activity and shows 50% or more of residual activity after a heat treatment at 45°C for 5 minutes.

11. A protein having the amino acid sequence of SEQ ID NO: 28 or the amino acid sequence of SEQ ID NO: 28 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues,

which has aconitase activity and shows 30% or more of residual activity after a heat treatment at 50°C for 3 minutes.

12. A protein having the amino acid sequence of SEQ ID NO: 30 or the amino acid sequence of SEQ ID NO: 30 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has isocitrate dehydrogenase activity and shows 50% or more of residual activity after a heat treatment at 45°C for 10 minutes.

13. A protein having the amino acid sequence of SEQ ID NO: 32 or the amino acid sequence of SEQ ID NO: 32 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has dihydrolipoamide dehydrogenase activity and is derived from *Corynebacterium thermoaminogenes*.

14. A protein having the amino acid sequence of SEQ ID NO: 34 or the amino acid sequence of SEQ ID NO: 34 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which has 2-oxoglutarate dehydrogenase activity and shows 30% or more of residual activity after a heat treatment at 50°C for 10 minutes.

15. A protein having the amino acid sequence of SEQ ID NO: 80 in Sequence Listing or the amino acid sequence of SEQ ID NO: 80 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which shows glutamate dehydrogenase activity at 42°C in an equivalent or higher degree compared with the activity at 37°C.

16. A protein having the amino acid sequence of SEQ ID NO: 90 in Sequence Listing or the amino acid sequence of SEQ ID NO: 90 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, which shows citrate synthase activity at 37°C in an equivalent or higher degree compared with the activity at 23°C.

17. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 2 or the amino acid sequence of SEQ ID NO: 2 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having isocitrate lyase activity.

18. The DNA according to Claim 17, which is a DNA defined in the following (a1) or (b1):

(a1) a DNA which comprises the nucleotide sequence of SEQ ID NO: 1 in Sequence Listing,

(b1) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 1 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having isocitrate lyase activity.

19. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 4 or the amino acid sequence of SEQ ID NO: 4 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and involved in acyl Co-A carboxylase activity.

20. The DNA according to Claim 19, which is a DNA defined in the following (a2) or (b2):

(a2) a DNA which comprises the nucleotide sequence of SEQ ID NO: 3 in Sequence Listing,

(b2) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 3 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein involved in acyl Co-A carboxylase activity.

21. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 6 or the amino acid sequence of SEQ ID NO: 6 including substitution, deletion, insertion, addition or inversion of one or

several amino acids residues, and having DtsR activity.

22. The DNA according to Claim 21, which is a DNA defined in the following (a3) or (b3):

(a3) a DNA which comprises the nucleotide sequence of SEQ ID NO: 5 in Sequence Listing,

(b3) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 5 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having DtsR activity.

23. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 8 or the amino acid sequence of SEQ ID NO: 8 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having DtsR activity.

24. The DNA according to Claim 23, which is a DNA defined in the following (a4) or (b4):

(a4) a DNA which comprises the nucleotide sequence of SEQ ID NO: 7 in Sequence Listing,

(b4) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 7 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having DtsR activity.

25. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 10 or the amino acid sequence of SEQ ID NO: 10 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having phosphofructokinase activity.

26. The DNA according to Claim 25, which is a DNA defined in the following (a5) or (b5):

(a5) a DNA which comprises the nucleotide sequence of SEQ ID NO: 9 in Sequence Listing,

(b5) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 9 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having phosphofructokinase activity.

27. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 93 or the amino acid sequence of SEQ ID NO: 93 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having invertase activity.

28. The DNA according to Claim 27, which is a DNA defined in the following (a6) or (b6):

(a6) a DNA which comprises the nucleotide sequence of SEQ ID NO: 93 in Sequence Listing,

(b6) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 93 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having invertase activity.

29. A DNA which codes for a protein having any one of the amino acid sequences of SEQ ID NOS: 17-20 or the amino acid sequence of any one of SEQ ID NOS: 17-20 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having a function involved in glutamic acid uptake.

30. The DNA according to Claim 29, which is a DNA defined in the following (a7) or (b7):

(a7) a DNA which comprises the nucleotide sequence of SEQ ID NO: 16 in Sequence Listing,

(b7) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 16 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having a function involved in glutamic acid uptake.

31. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 22 or the amino acid

sequence of SEQ ID NO: 22 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having pyruvate dehydrogenase activity.

32. The DNA according to Claim 31, which is a DNA defined in the following (a8) or (b8):

(a8) a DNA which comprises the nucleotide sequence of SEQ ID NO: 21 in Sequence Listing,

(b8) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 21 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having pyruvate dehydrogenase activity.

33. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 24 or the amino acid sequence of SEQ ID NO: 24 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having pyruvate carboxylase activity.

34. A DNA according to Claim 33, which is a DNA defined in the following (a9) or (b9):

(a9) a DNA which comprises the nucleotide sequence of SEQ ID NO: 23 in Sequence Listing,

(b9) a DNA which is hybridizable with the



nucleotide sequence of SEQ ID NO: 23 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having pyruvate carboxylase activity.

35. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 26 or the amino acid sequence of SEQ ID NO: 26 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having phosphoenolpyruvate carboxylase activity.

36. The DNA according to Claim 35, which is a DNA defined in the following (a10) or (b10):

(a10) a DNA which comprises the nucleotide sequence of SEQ ID NO: 25 in Sequence Listing,

(b10) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 25 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having phosphoenolpyruvate carboxylase activity.

37. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 28 or the amino acid sequence of SEQ ID NO: 28 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having aconitase

activity.

38. The DNA according to Claim 37, which is a DNA defined in the following (a11) or (b11):

(a11) a DNA which comprises the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing,

(b11) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having aconitase activity.

39. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 30 or the amino acid sequence of SEQ ID NO: 30 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having isocitrate dehydrogenase activity.

40. The DNA according to Claim 39, which is a DNA defined in the following (a12) or (b12):

(a12) a DNA which comprises the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing,

(b12) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 27 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein

having isocitrate dehydrogenase activity.

41. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 32 or the amino acid sequence of SEQ ID NO: 32 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having dihydrolipoamide dehydrogenase activity.

42. The DNA according to Claim 41, which is a DNA defined in the following (a13) or (b13):

(a13) a DNA which comprises the nucleotide sequence of SEQ ID NO: 31 in Sequence Listing,

(b13) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 31 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having dihydrolipoamide dehydrogenase activity.

43. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 34 or the amino acid sequence of SEQ ID NO: 34 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and having 2-oxoglutarate dehydrogenase activity.

44. The DNA according to Claim 43, which is a

DNA defined in the following (a14) or (b14):

(a14) a DNA which comprises the nucleotide sequence of SEQ ID NO: 33 in Sequence Listing,

(b14) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 33 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein having 2-oxoglutarate dehydrogenase activity.

45. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 80 in Sequence Listing or the amino acid sequence of SEQ ID NO: 80 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and showing glutamate dehydrogenase activity at 42°C in an equivalent or higher degree compared with the activity at 37°C.

46. The DNA according to Claim 45, which is a DNA defined in the following (a15) or (b15):

(a15) a DNA which comprises the nucleotide sequence of SEQ ID NO: 79 in Sequence Listing,

(b15) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 79 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein showing glutamate dehydrogenase activity at 42°C in an

equivalent or higher degree compared with the activity at 37° C.

47. A DNA which codes for a protein having the amino acid sequence of SEQ ID NO: 90 in Sequence Listing or the amino acid sequence of SEQ ID NO: 90 including substitution, deletion, insertion, addition or inversion of one or several amino acids residues, and showing citrate synthase activity at 37° C in an equivalent or higher degree compared with the activity at 23° C.

48. The DNA according to Claims 47, which is a DNA defined in the following (a16) or (b16):

(a16) a DNA which comprises the nucleotide sequence of SEQ ID NO: 89 in Sequence Listing,

(b16) a DNA which is hybridizable with the nucleotide sequence of SEQ ID NO: 89 in Sequence Listing or a primer prepared based on the nucleotide sequence under a stringent condition, and codes for a protein showing citrate synthase activity at 37° C in an equivalent or higher degree compared with the activity at 23° C.

49. A method for producing L-amino acid, which comprises culturing a microorganism introduced with a DNA according to any one of Claims 17 to 48 in a medium to produce and accumulate L-amino acid in the medium,

and collecting the L-amino acid from the medium.

## ABSTRACT

A plurality of primer sets are designed based on a region where conservation at the amino acid level is observed among various microorganisms for known gene sequences corresponding to a gene coding for an enzyme of the L-amino acid biosynthetic pathway derived from *Corynebacterium thermoaminogenes*, preferably an enzyme that functions at a higher temperature compared with that of *Corynebacterium glutamicum*. PCR is performed by using the primers and chromosomal DNA of *Corynebacterium thermoaminogenes* as a template. The primers with which an amplification fragment has been obtained are used as primers for screening to select a clone containing a target DNA fragment from a plasmid library of chromosomal DNA of *Corynebacterium thermoaminogenes*.

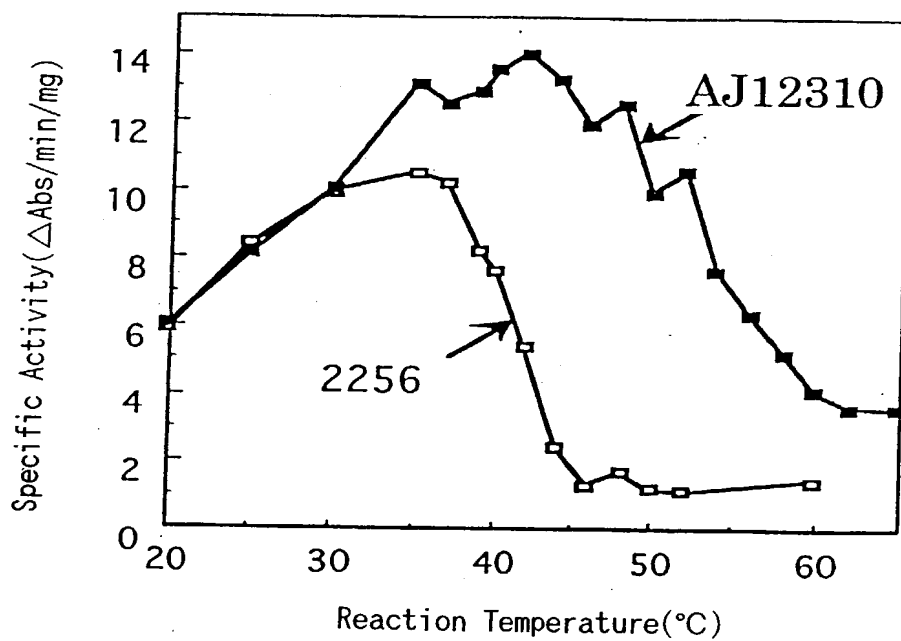


Fig. 1

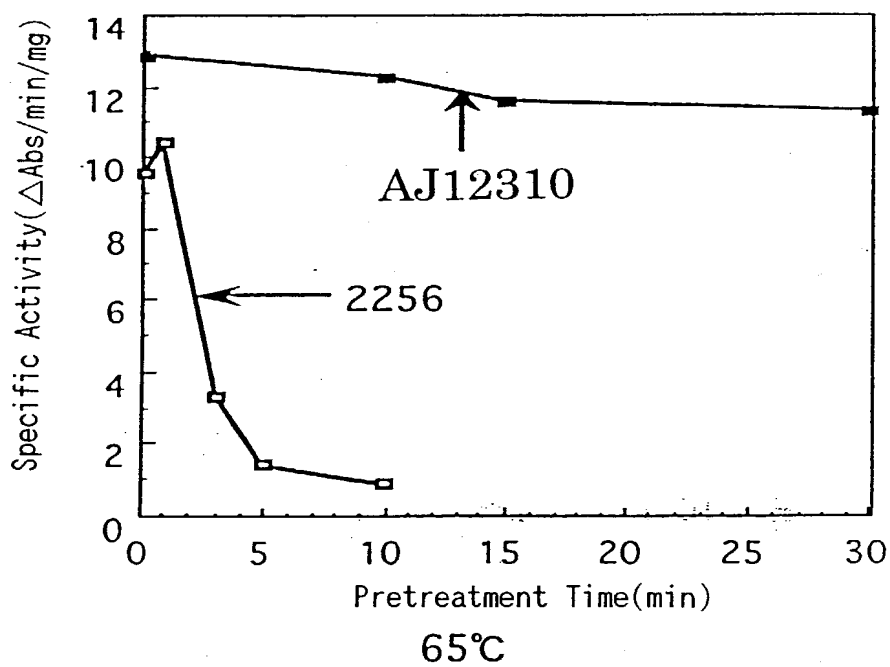


Fig. 2



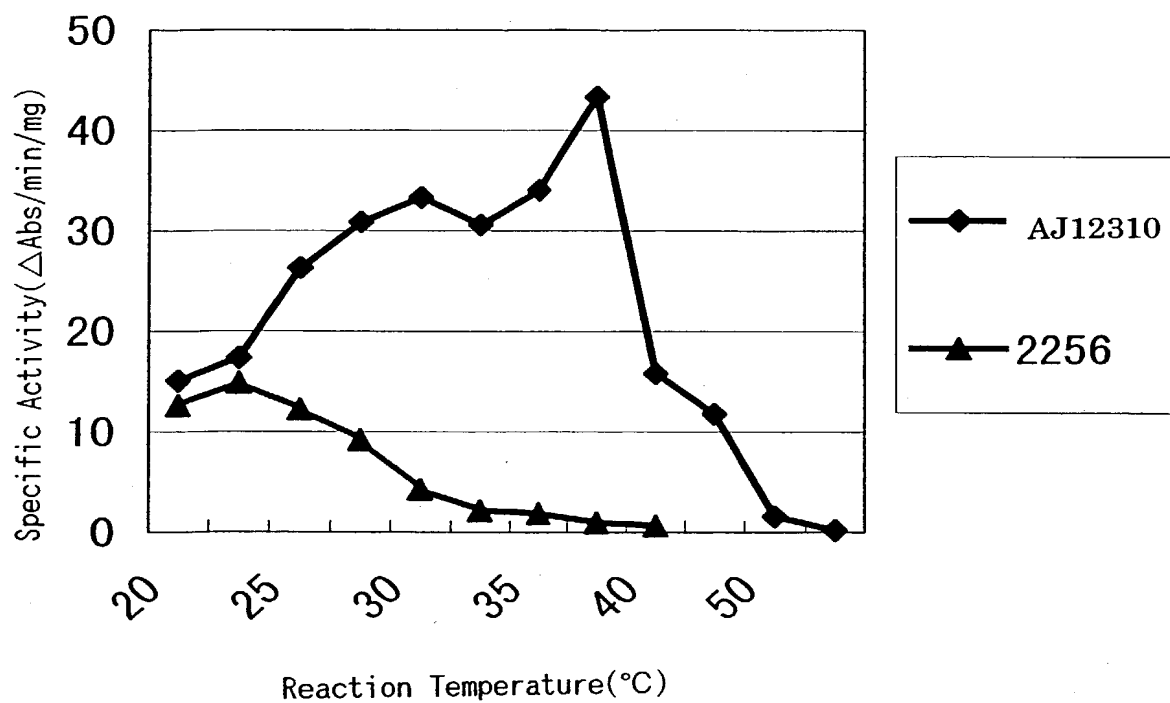


Fig. 3

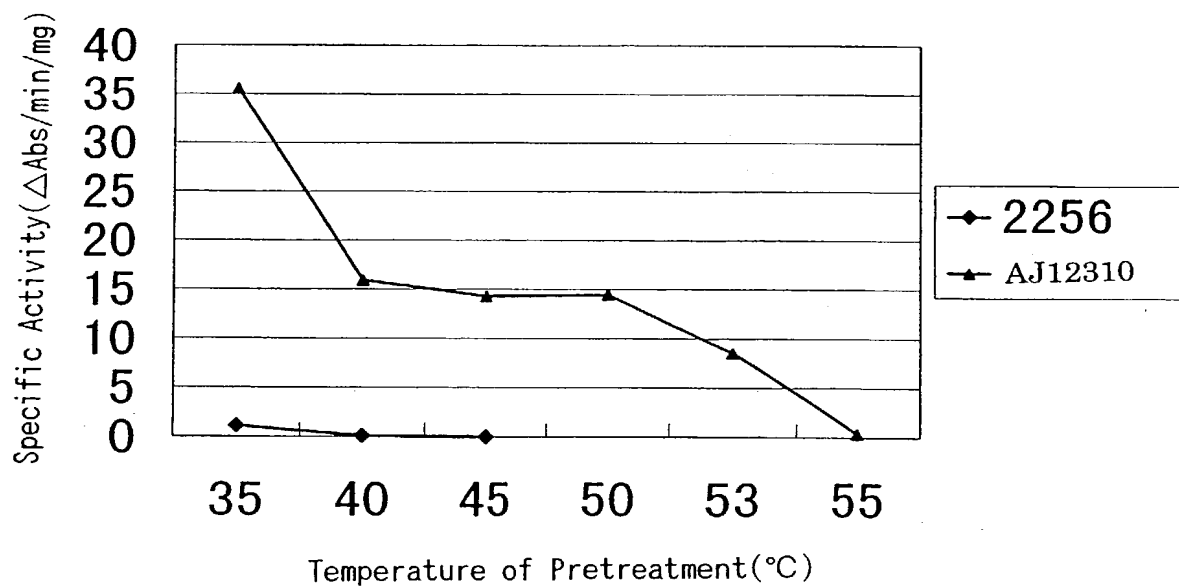


Fig. 4

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Fig. 5

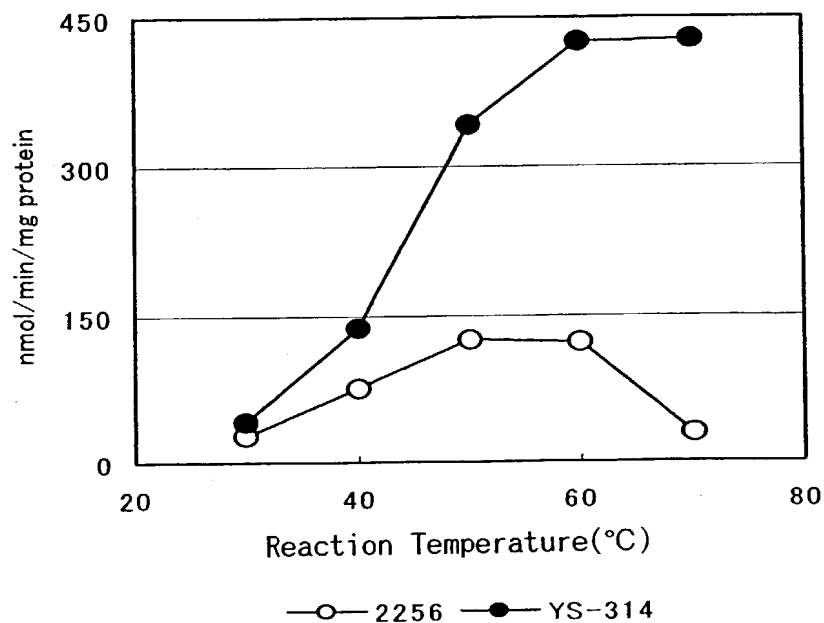


Fig. 6

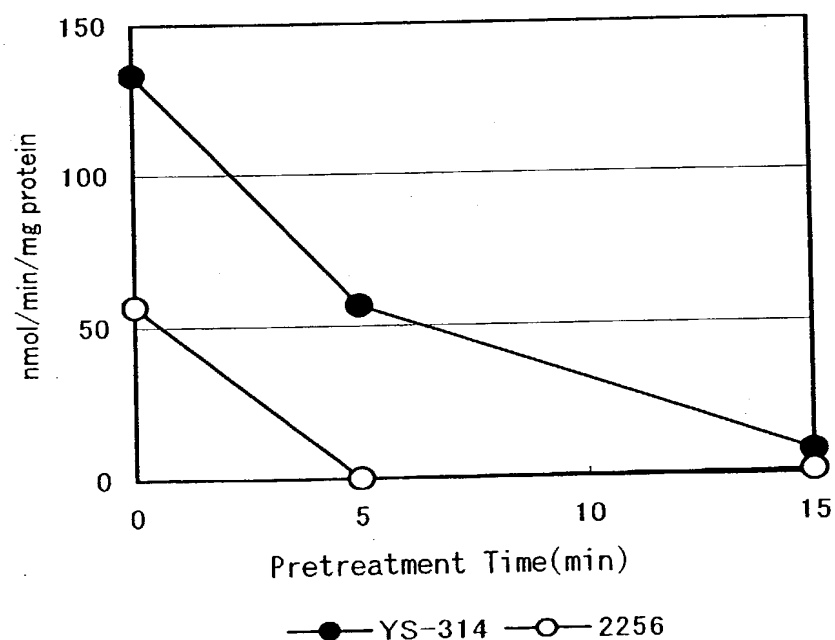


Fig. 7

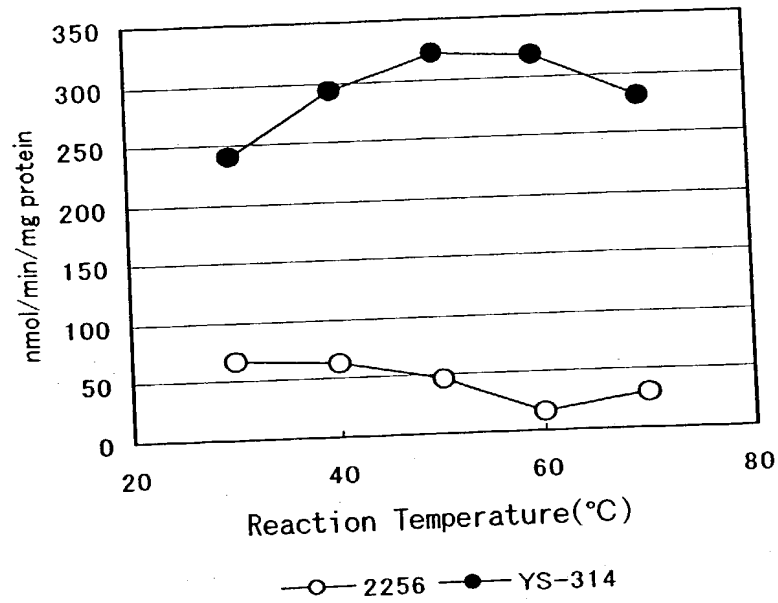
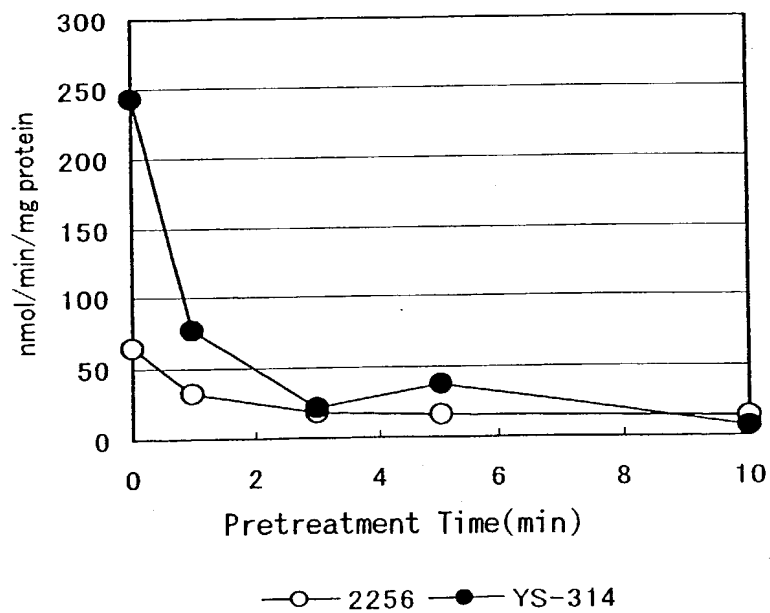


Fig. 8



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Fig. 9

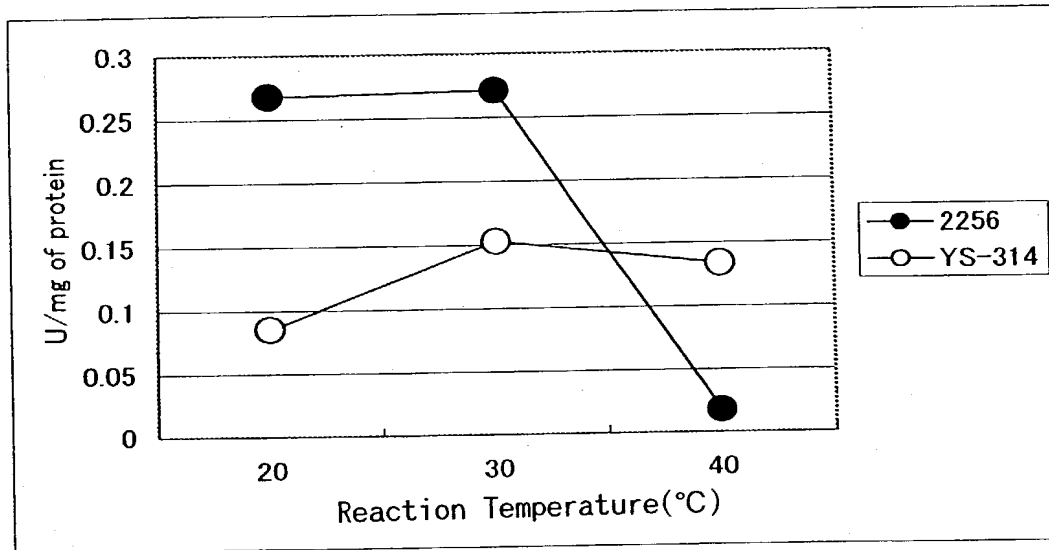


Fig. 10

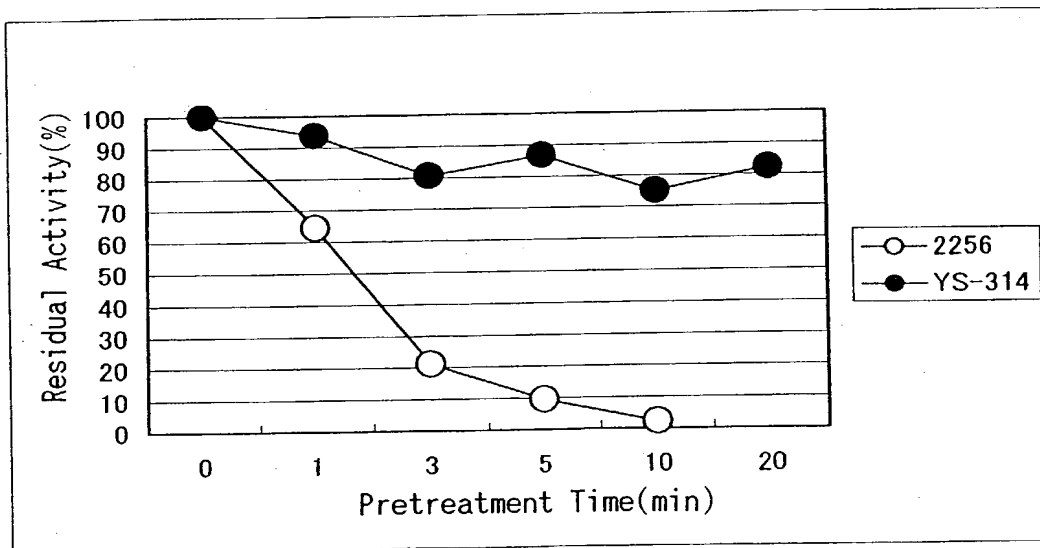


Fig. 11

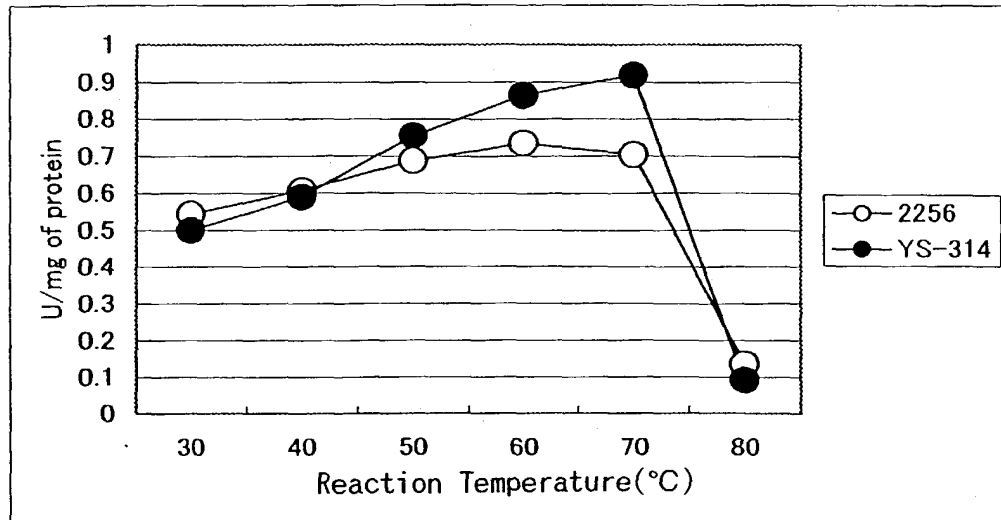


Fig. 12

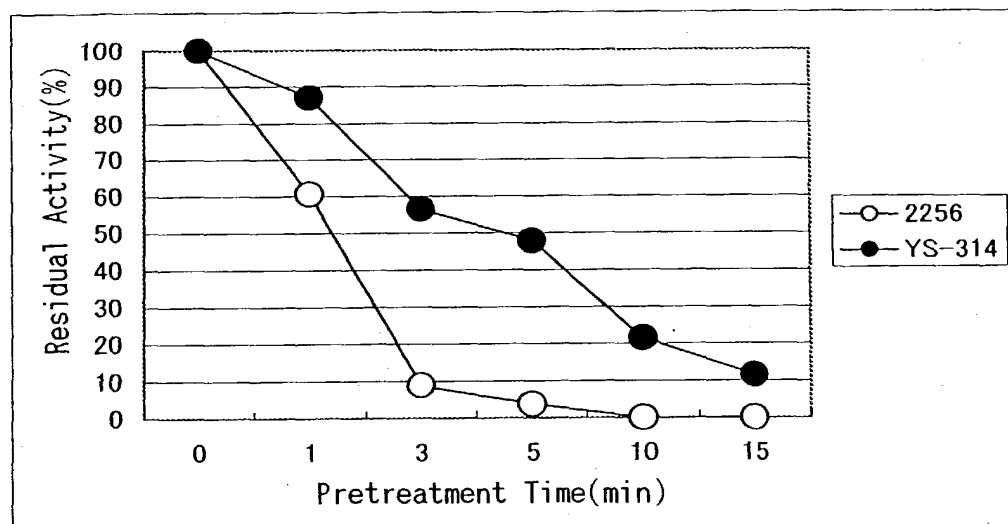


Fig. 13

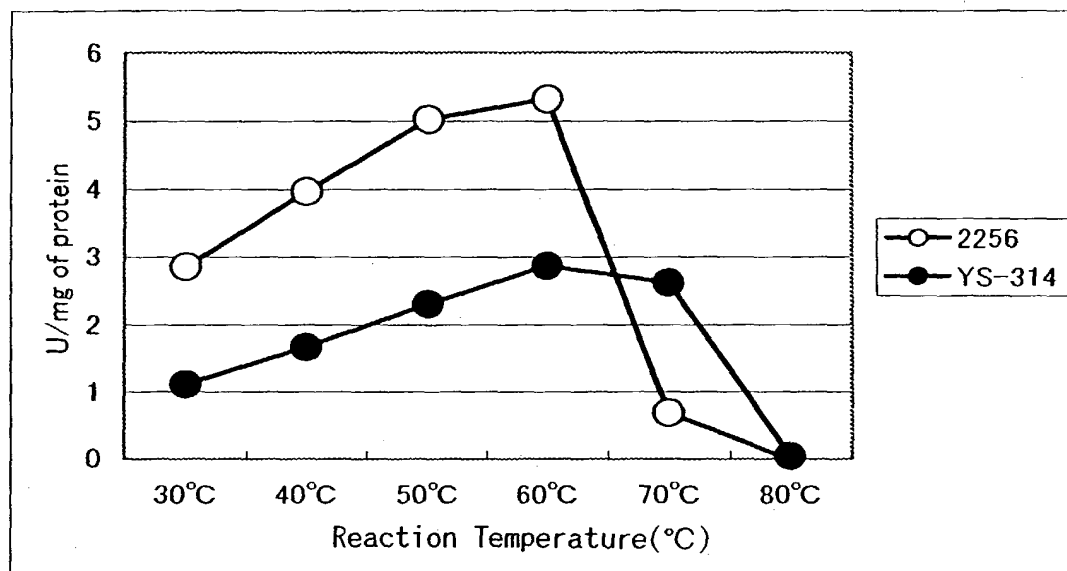
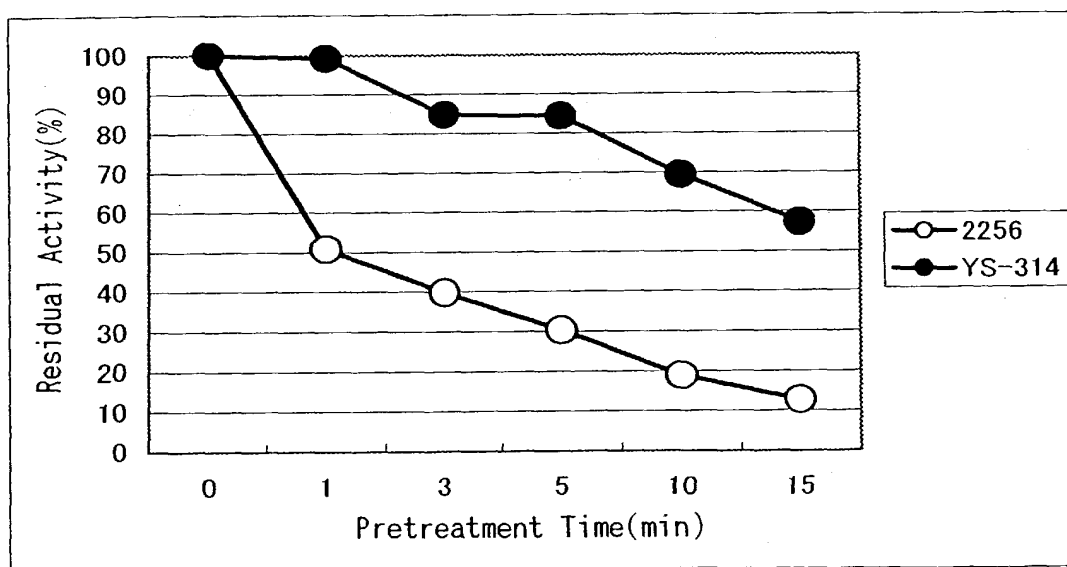
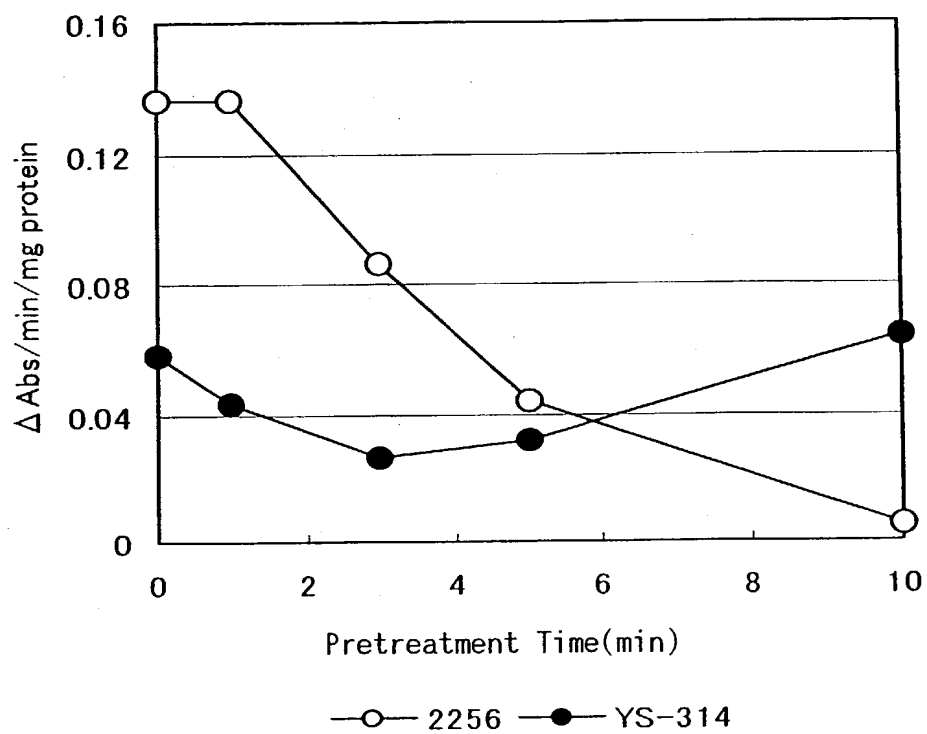


Fig. 14



*Fig. 15*

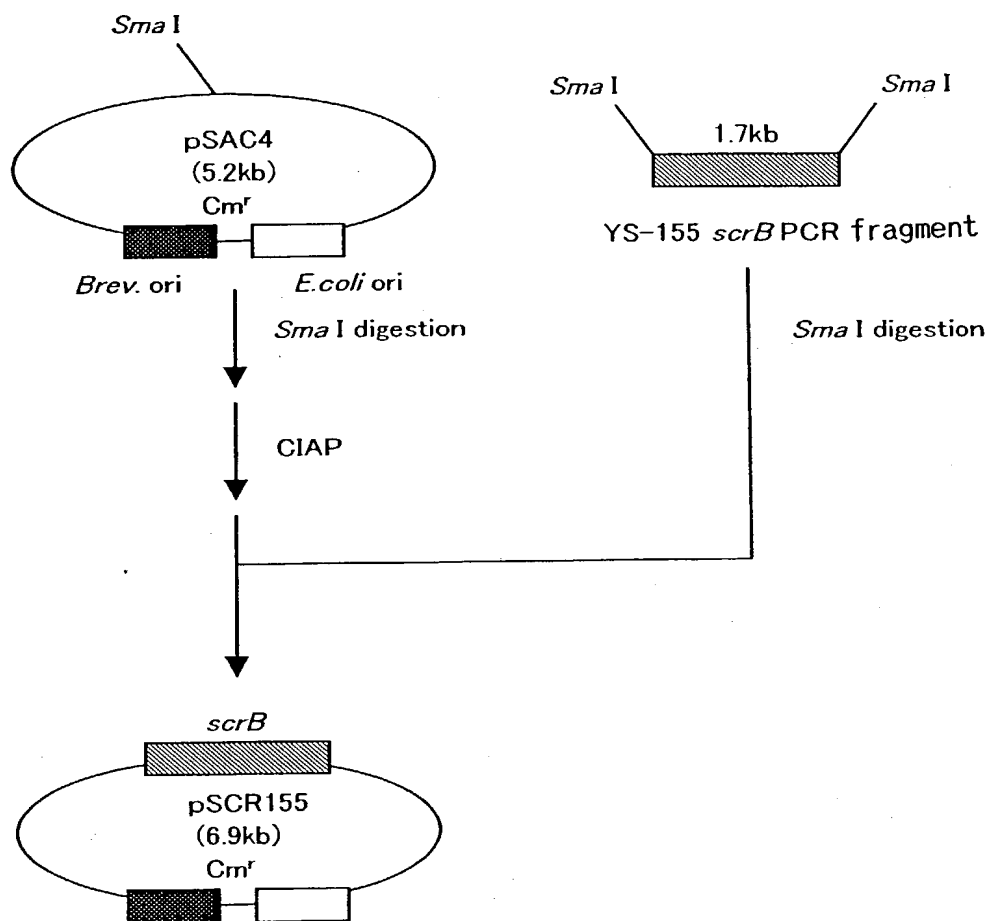


Fig. 16



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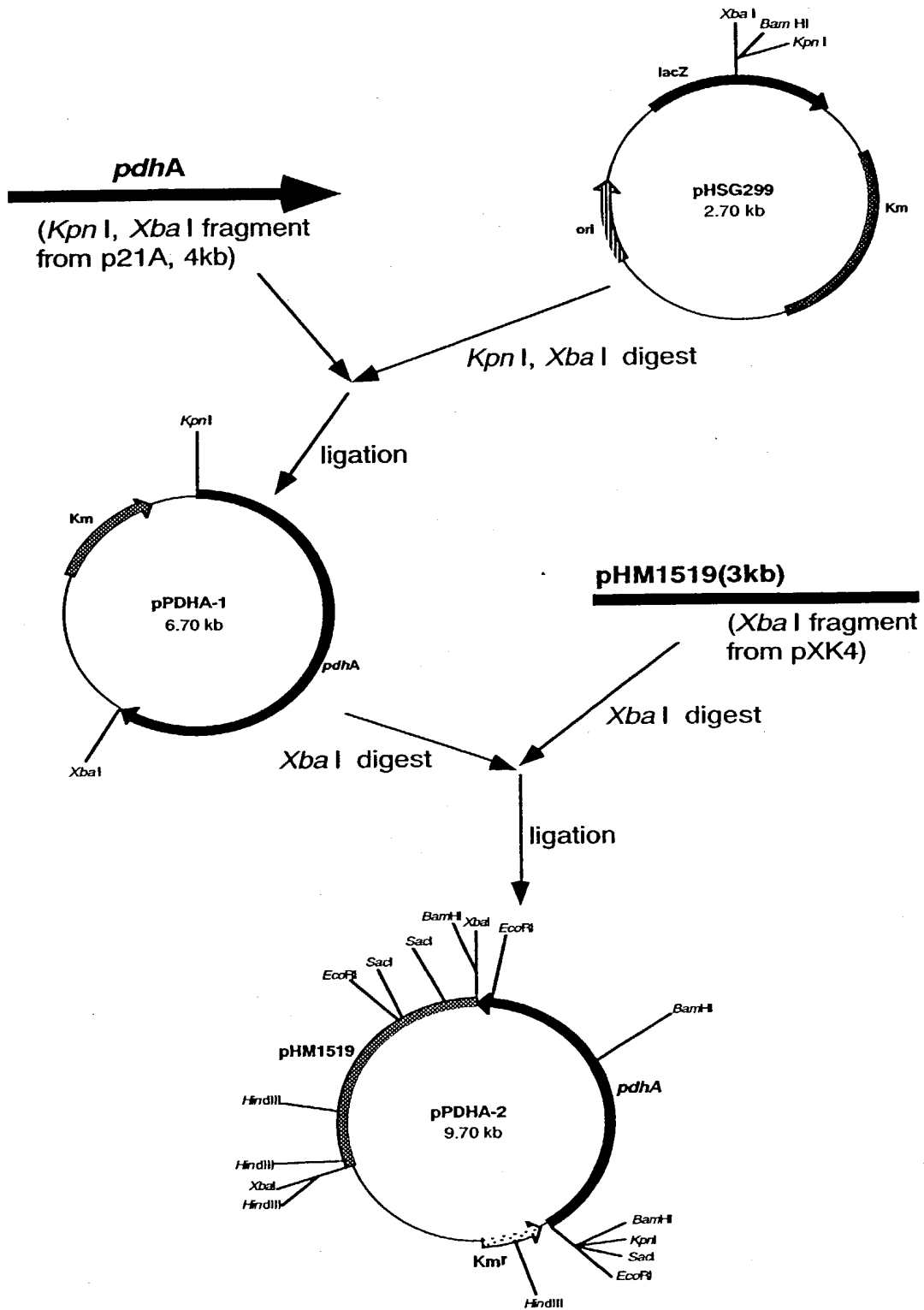


Fig. 17

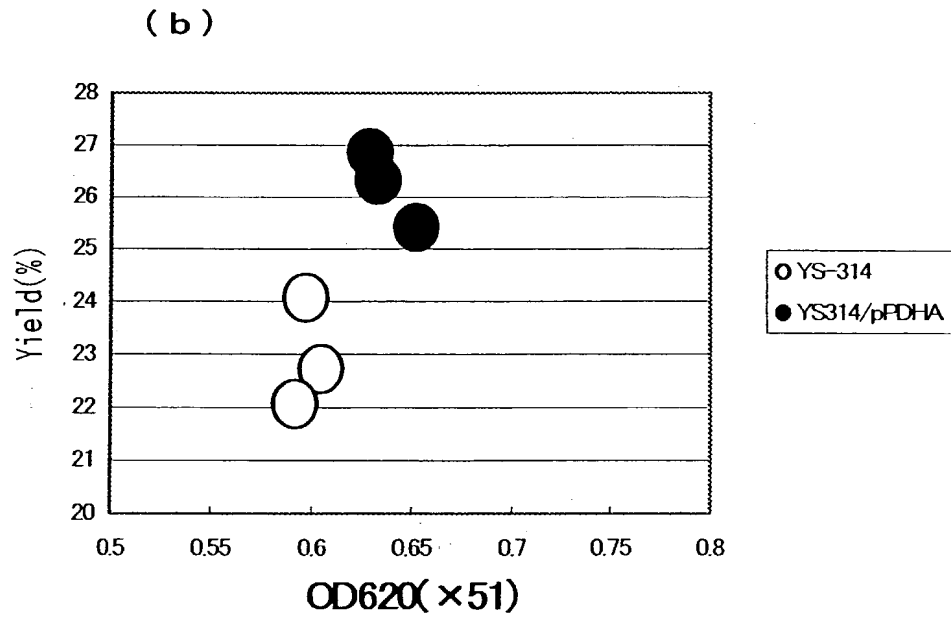
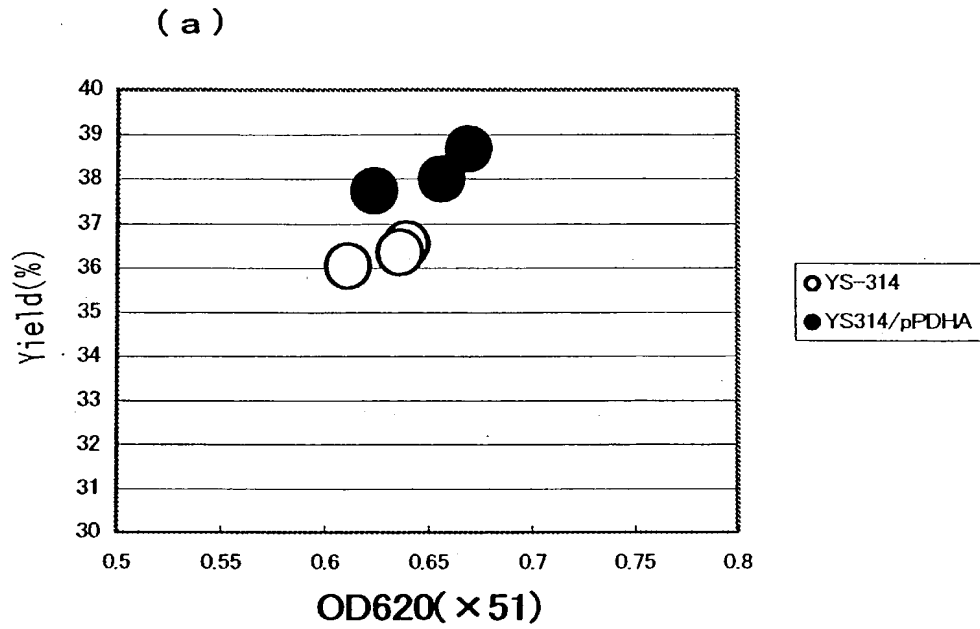


Fig. 18

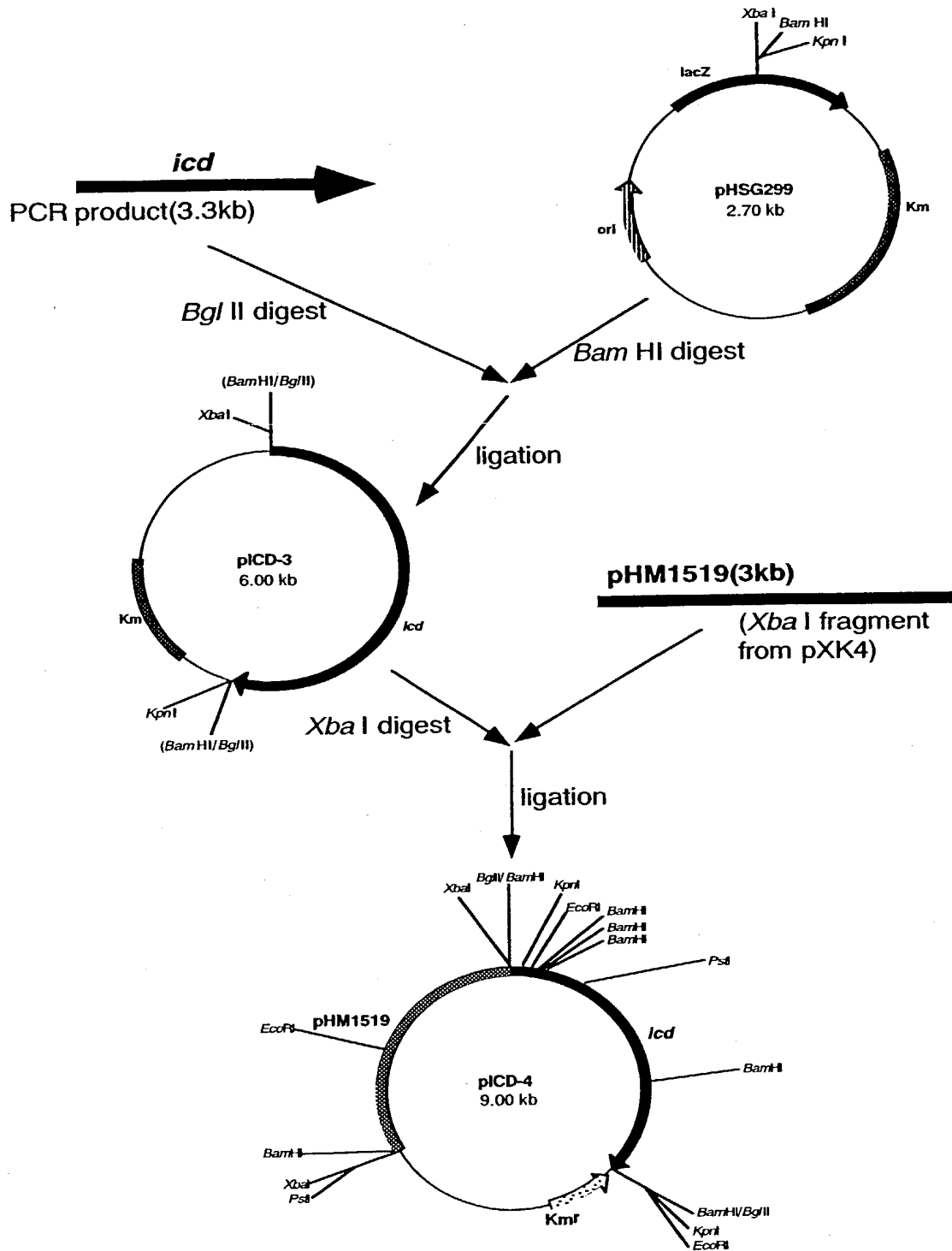


Fig. 19

OBLON ET AL (703) 413-3000

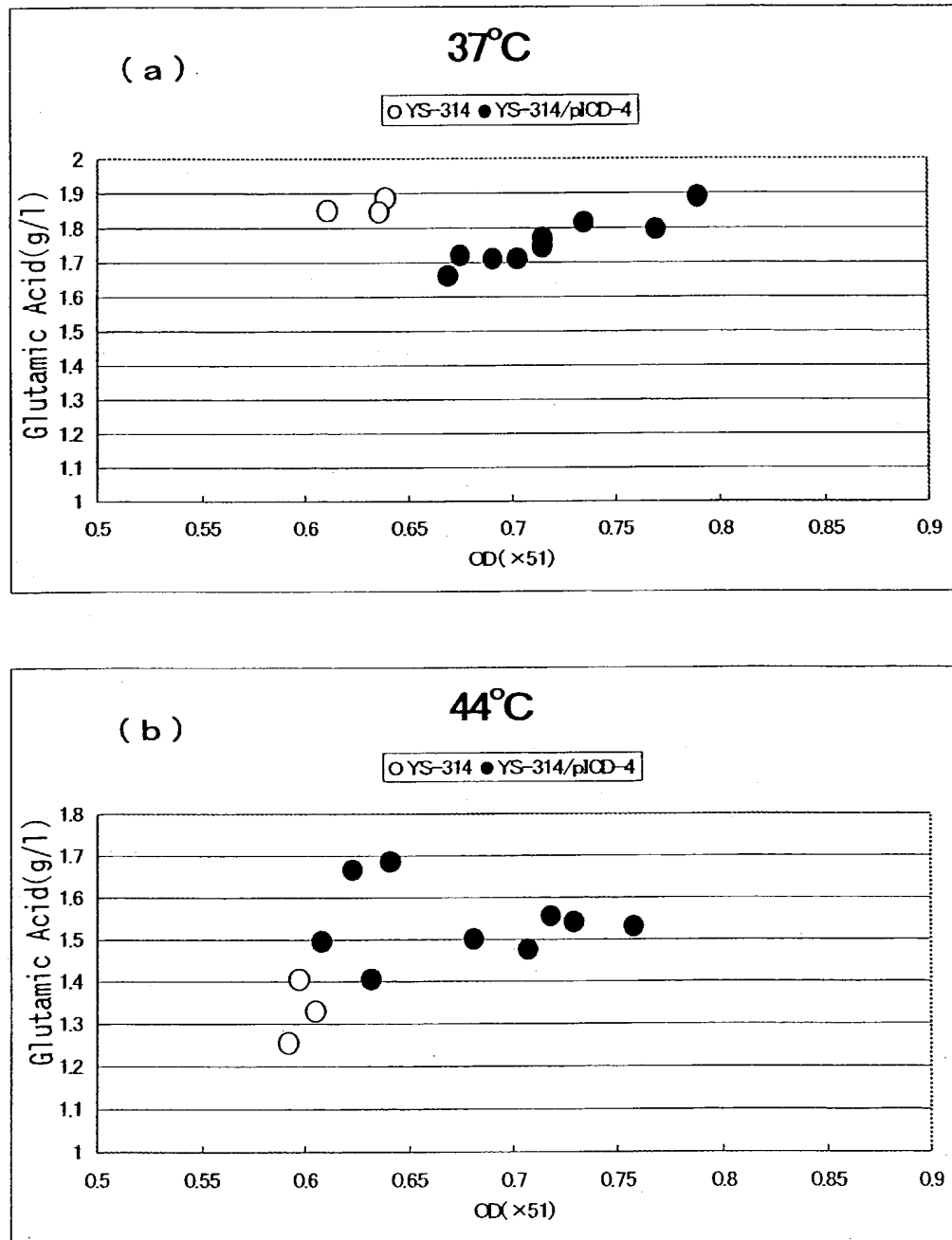
DOCKET # 22159US SHEET 13 OF 15

Fig. 20

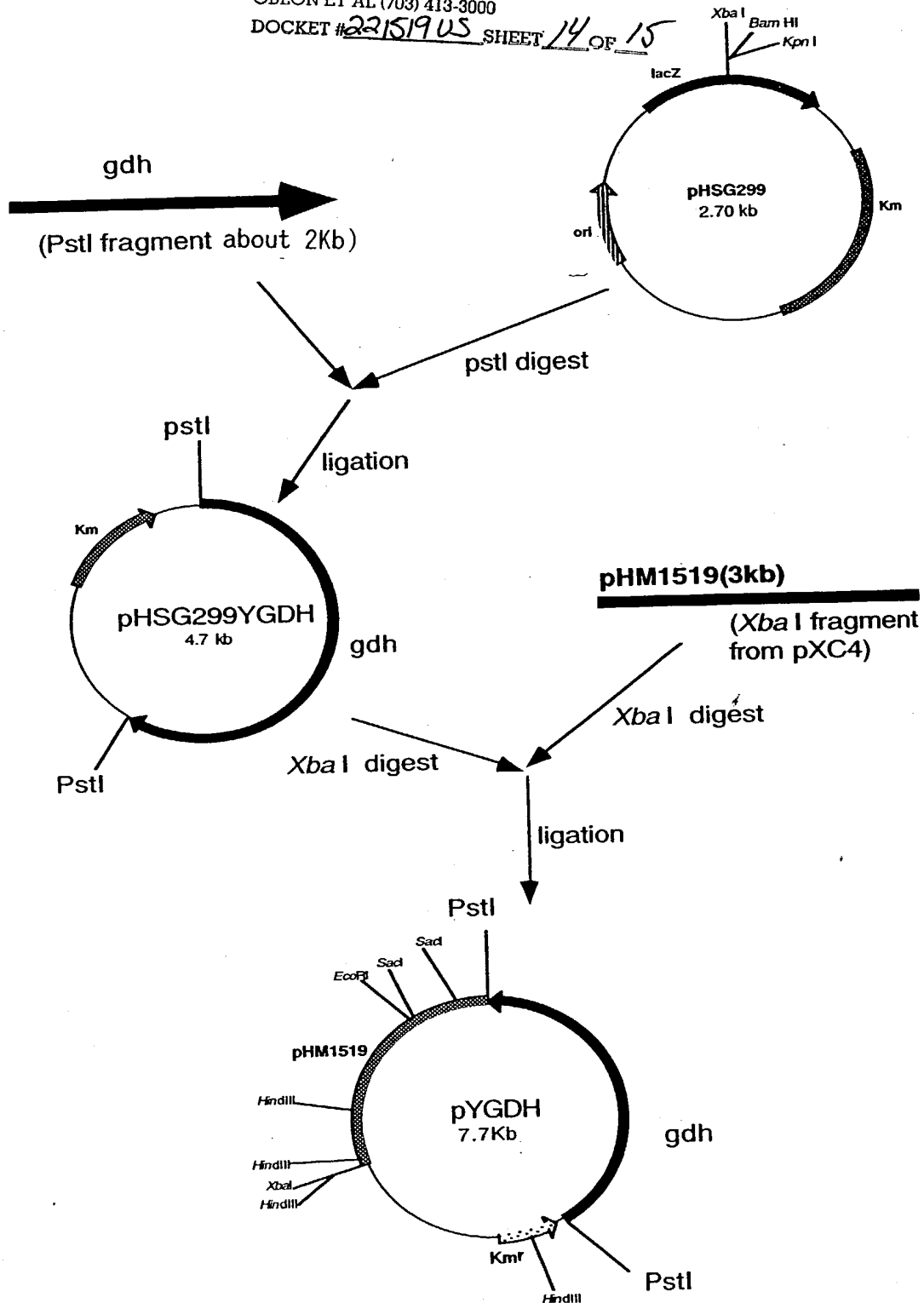


Fig. 21

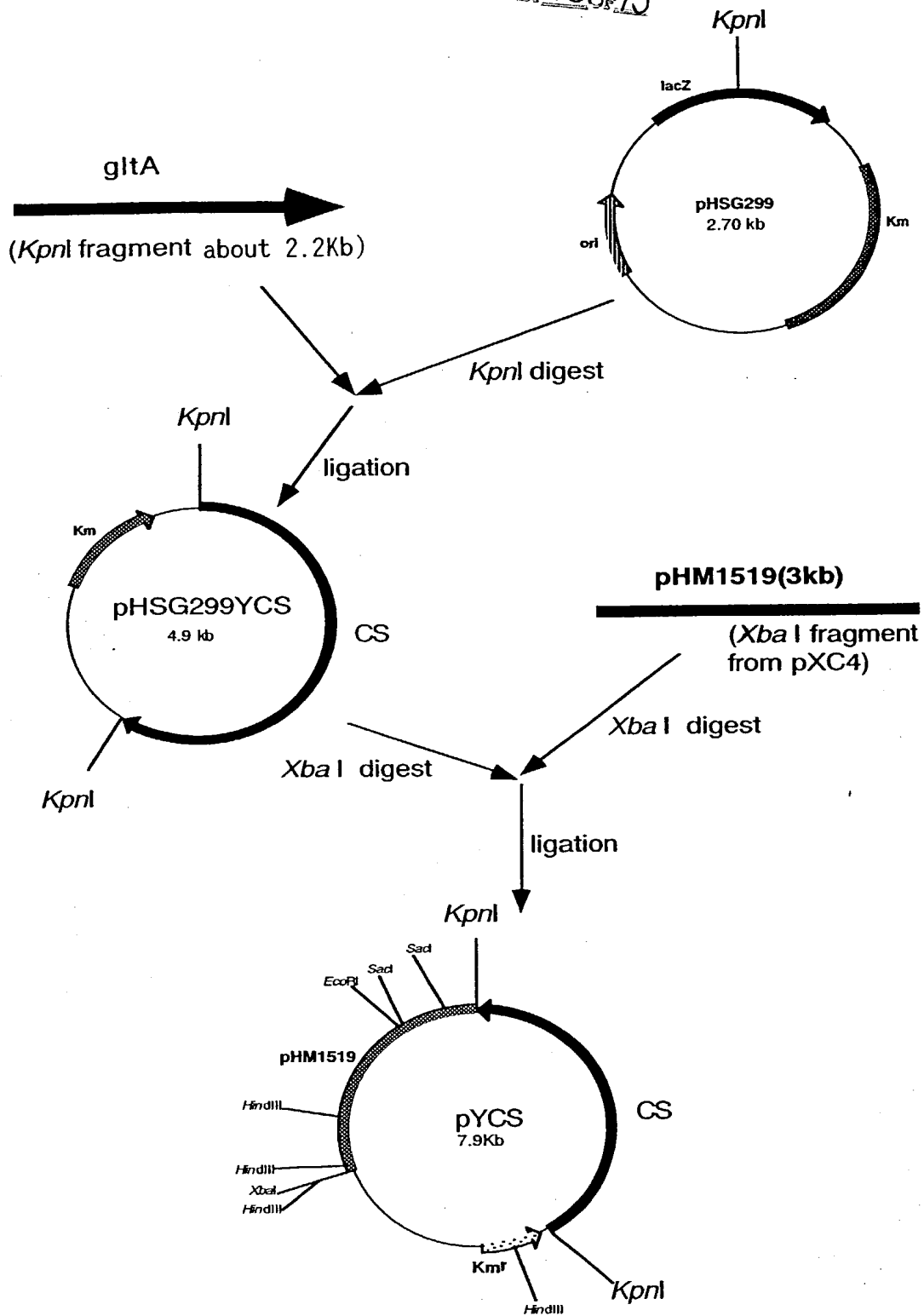


Fig. 22

# Declaration, Power Of Attorney and Petition

Page 1 of 5

WE (I) the undersigned inventor(s), hereby declare(s) that:

My residence, post office address and citizenship are as stated below next to my name,

We (I) believe that we are (I am) the original, first, and joint (sole) inventor(s) of the subject matter which is claimed and for which a patent is sought on the invention entitled

GENES FOR HEAT RESISTANT ENZYMES OF AMINO ACID BIOSYNTHETIC PATHWAY  
DERIVED FROM THERMOPHILIC CORYNEFORM BACTERIA

the specification of which

- ☐ is attached hereto.
- ☐ was filed on \_\_\_\_\_ as  
 Application Serial No. \_\_\_\_\_  
 and amended on \_\_\_\_\_.
- ☒ was filed as PCT international application  
 Number PCT/JP 00/06913  
 on October 4, 2000,  
 and was amended under PCT Article 19  
 on \_\_\_\_\_ (if applicable).

We (I) hereby state that we (I) have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We (I) acknowledge the duty to disclose information known to be material to the patentability of this application as defined in Section 1.56 of Title 37 Code of Federal Regulations.

We (I) hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s)

Application No.	Country	Day/Month/Year	Priority Claimed
<u>11-282716</u>	<u>Japan</u>	<u>04/10/1999</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
<u>11-311147</u>	<u>Japan</u>	<u>01/11/1999</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
<u>2000-120687</u>	<u>Japan</u>	<u>21/04/2000</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
_____	_____	_____	<input type="checkbox"/> Yes <input type="checkbox"/> No

We (I) hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below.

(Application Number)

(Filing Date)

(Application Number)

(Filing Date)

We (I) hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.

Filing Date

Status (pending, patented,  
abandoned)

Application Serial No.	Filing Date	Status (pending, patented, abandoned)

24 And we (I) hereby appoint: Norman F. Oblon, Registration Number 24,618; Marvin J. Spivak, Registration Number 24,913; C. Irvin McClelland, Registration Number 21,124; Gregory J. Maier, Registration Number 25,599; Arthur I. Neustadt, Registration Number 24,854; Richard D. Kelly, Registration Number 27,757; James D. Hamilton, Registration Number 28,421; Eckhard H. Kuesters, Registration Number 28,870; Robert T. Pous, Registration Number 29,099; Charles L. Gholz, Registration Number 26,395; Vincent J. Sunderdick, Registration Number 29,004; William E. Beaumont, Registration Number 30,996; Steven B. Kelber, Registration Number 30,073; Robert F. Gnuse, Registration Number 27,295; Jean-Paul Lavalleye, Registration Number 31,451; Timothy R. Schwartz, Registration Number 32,171; Stephen G. Baxter, Registration Number 32,884; Martin M. Zoltick, Registration Number 35,745; Robert W. Hahl, Registration Number 33,893; Richard L. Treanor, Registration Number 36,379; Steven P. Weihrouch, Registration Number 32,829; John T. Goolkasian, Registration Number 26,142; Marc R. Labgold, Registration Number 34,651; William J. Healey, Registration Number 36,160; and Richard L. Chinn, Registration Number 34,305; our (my) attorneys, with full powers of substitution and revocation, to prosecute this application and to transact all business in the Patent Office connected therewith; and we (I) hereby request that all correspondence regarding this application be sent to the firm of OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT, P.C., whose Post Office Address is: Fourth Floor, 1755 Jefferson Davis Highway, Arlington, Virginia 22202.

We (I) declare that all statements made herein of our (my) own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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Signature of Inventor

Date

1/123

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&lt;110&gt; Ajinomoto Co., Inc.

<120> Genes for Heat resistant Enzymes of Amino Acid  
Biosynthetic Pathway Derived from Thermophilic  
Coryneform Bacteria

&lt;130&gt; OP1072B691SM

&lt;140&gt;

&lt;141&gt; 2000-10-04

&lt;150&gt; JP 11-282716

&lt;151&gt; 1999-10-04

&lt;150&gt; JP 11-311147

&lt;151&gt; 1999-11-01

&lt;150&gt; JP 2000-120687

&lt;151&gt; 2000-04-21

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2/123

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Pro Arg Thr Ala Gln Glu Ile Gln Gln Asp Trp Asp Thr Asn Pro Arg						
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tgg aac gga atc acc cgc gac tac acc gct gag cag gta gct gag ctc						690
Trp Asn Gly Ile Thr Arg Asp Tyr Thr Ala Glu Gln Val Ala Glu Leu						
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Gln Gly Ser Val Val Glu Glu His Thr Leu Ala Lys Arg Gly Ala Glu						
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Ile Leu Trp Asp Ala Val Ser Ala Glu Gly Asp Asp Tyr Ile Asn Ala						
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Leu Gly Ala Leu Thr Gly Asn Gln Ala Val Gln Gln Val Arg Ala Gly						
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Leu Lys Ala Val Tyr Leu Ser Gly Trp Gln Val Ala Gly Asp Ala Asn						
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Leu Ala Gly His Thr Tyr Pro Asp Gln Ser Leu Tyr Pro Ala Asn Ser						
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gtc ccg aac gtt gtc cgt cgc atc aac aac gca ctg ctg cgc gcc gat						978
Val Pro Asn Val Val Arg Arg Ile Asn Asn Ala Leu Leu Arg Ala Asp						
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Glu Ile Ala Arg Val Glu Gly Asp Thr Ser Val Asp Asn Trp Leu Val						
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3/123

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Phe	Val	Asp	Leu	Gln	Asn	Arg	Glu	Phe	Lys	Ala	Ala	Glu	Glu	Arg	Gly	
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ttc	acc	gcc	gtc	aag	cac	cag	cgt	gag	gtc	ggc	gcc	ggc	tac	ttc	gac	1794
Phe	Thr	Ala	Val	Lys	His	Gln	Arg	Glu	Val	Gly	Ala	Gly	Tyr	Phe	Asp	
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4/123

425

430

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&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

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Gly	Ala	Ala	Gly	Thr	His	Trp	Glu	Asp	Gln	Leu	Ala	Ser	Glu	Lys	Lys
		180					185						190		
Cys	Gly	His	Leu	Gly	Gly	Lys	Val	Leu	Ile	Pro	Thr	Gln	Gln	His	Ile
		195				200						205			
Arg	Thr	Leu	Asn	Ser	Ala	Arg	Leu	Ala	Ala	Asp	Val	Ala	Asn	Thr	Pro
		210				215					220				
Thr	Val	Val	Ile	Ala	Arg	Thr	Asp	Ala	Glu	Ala	Ala	Thr	Leu	Ile	Thr
225				230					235					240	
Ser	Asp	Val	Asp	Glu	Arg	Asp	Arg	Pro	Phe	Ile	Thr	Gly	Glu	Arg	Thr
			245					250					255		
Ala	Glu	Gly	Tyr	Tyr	His	Val	Lys	Pro	Gly	Leu	Glu	Pro	Cys	Ile	Ala

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	260		265		270										
Arg	Ala	Lys	Ser	Tyr	Ala	Pro	Tyr	Ala	Asp	Met	Ile	Trp	Met	Glu	Thr
	275						280						285		
Gly	Thr	Pro	Asp	Leu	Glu	Leu	Ala	Lys	Lys	Phe	Ala	Glu	Gly	Val	Arg
	290						295						300		
Ser	Glu	Phe	Pro	Asp	Gln	Leu	Leu	Ser	Tyr	Asn	Cys	Ser	Pro	Ser	Phe
305					310						315				320
Asn	Trp	Ser	Ala	His	Leu	Glu	Ala	Asp	Glu	Ile	Ala	Lys	Phe	Gln	Lys
				325						330				335	
Glu	Leu	Gly	Ala	Met	Gly	Phe	Lys	Phe	Gln	Phe	Ile	Thr	Leu	Ala	Gly
		340						345					350		
Phe	His	Ser	Leu	Asn	Tyr	Gly	Met	Phe	Asp	Leu	Ala	Tyr	Gly	Tyr	Ala
	355						360						365		
Arg	Glu	Gly	Met	Pro	Ala	Phe	Val	Asp	Leu	Gln	Asn	Arg	Glu	Phe	Lys
	370						375						380		
Ala	Ala	Glu	Glu	Arg	Gly	Phe	Thr	Ala	Val	Lys	His	Gln	Arg	Glu	Val
385					390					395					400
Gly	Ala	Gly	Tyr	Phe	Asp	Thr	Ile	Ala	Thr	Thr	Val	Asp	Pro	Asn	Ser
				405					410					415	
Ser	Thr	Thr	Ala	Leu	Lys	Gly	Ser	Thr	Glu	Glu	Cys	Gln	Phe	His	
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&lt;210&gt; 3

&lt;211&gt; 2381

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (577)..(2349)

&lt;400&gt; 3

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cacctagtc accccggggt caccctccig gtcacccccg taccctcccg ggtacacccc 300
ggggacgggg tigtaccigg atctccctcg catgtggaca ccgggaaact ttgacctggga 360
aalgacatc cagtaccgta atgcgggtat gttaacgcgg tcacagggtta caccagaatc 420
cggatcgtct aaccccccta gcgggatcgc ctaaaagatc accgagttag tigtcaagaa 480
taatgcgat cgcaggggca ctgcatatcg ctgtcatgca gtcaatgaac agtgcgggtc 540
ctgtcgtga agaaaatcaa aaccaggagg gtttta gtc tca gtc gag acc agg 594
Val Ser Val Glu Thr Arg

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aag atc acc aag gta ctt gtc gcc aac cgt ggt gaa atc gca atc cgt	1	5	642
Lys Ile Thr Lys Val Leu Val Ala Asn Arg Gly Glu Ile Ala Ile Arg			
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gtt ttc cgc gca gca cgg gat gaa ggc atc gcc tct gtc gcc gtc tac			690
Val Phe Arg Ala Ala Arg Asp Glu Gly Ile Ala Ser Val Ala Val Tyr			
25	30	35	
gcg gag ccg gac gca gat gcc cct ttc gtc gag tat gcc gat gag gcc			738
Ala Glu Pro Asp Ala Asp Ala Pro Phe Val Glu Tyr Ala Asp Glu Ala			
40	45	50	
ttc gca ctc ggt ggc cag act tcc gca gag tcc tac ctc gtc att gac			786
Phe Ala Leu Gly Gly Gln Thr Ser Ala Glu Ser Tyr Leu Val Ile Asp			
55	60	65	70
aag atc att gac gca gca cgc aag tcc ggt gca gac gct gtc cac ccc			834
Lys Ile Ile Asp Ala Ala Arg Lys Ser Gly Ala Asp Ala Val His Pro			
75	80	85	
ggc tac ggc ttc ctc gcc gag aac gcc gat ttc gct gaa gct gtc atc			882
Gly Tyr Gly Phe Leu Ala Glu Asn Ala Asp Phe Ala Glu Ala Val Ile			
90	95	100	
aac gag ggc ctg atc tgg atc gga cca tcc cct gag tcc atc cgt tcc			930
Asn Glu Gly Leu Ile Trp Ile Gly Pro Ser Pro Glu Ser Ile Arg Ser			
105	110	115	
ctc ggt gac aag gtc acc gca cgc cac atc gcc aac aac gcc aac gca			978
Leu Gly Asp Lys Val Thr Ala Arg His Ile Ala Asn Asn Ala Asn Ala			
120	125	130	
ccg atg gca ccg ggc acc aag gag cct gtc aag gac gcc gct gag gtt			1026
Pro Met Ala Pro Gly Thr Lys Glu Pro Val Lys Asp Ala Ala Glu Val			
135	140	145	150
gtc gcc ttc gcc gag gag ttc ggt ctc ccc atc gcc atc aag gct gcc			1074
Val Ala Phe Ala Glu Glu Phe Gly Leu Pro Ile Ala Ile Lys Ala Ala			
155	160	165	
ttc ggt ggc ggc gga cgt ggc atg aag gtc gcc tac gag atg gac gag			1122
Phe Gly Gly Gly Gly Arg Gly Met Lys Val Ala Tyr Glu Met Asp Glu			
170	175	180	
gtc gcc gac ctc ttc gaa tcc gcc acc cgt gag gcc acc gcc gcc ttc			1170
Val Ala Asp Leu Phe Glu Ser Ala Thr Arg Glu Ala Thr Ala Ala Phe			
185	190	195	
ggt cgt ggt gag tgc ttc gtg gag cgc tac ctg gac aag gcc cgc cac			1218
Gly Arg Gly Glu Cys Phe Val Glu Arg Tyr Leu Asp Lys Ala Arg His			
200	205	210	
gtc gag gca cag gtc atc gcc gac aag cac ggc aac gtt gtg gtc gcc			1266
Val Glu Ala Gln Val Ile Ala Asp Lys His Gly Asn Val Val Val Ala			
215	220	225	230

ggt acc cgt gac tgc tcc ctg cag cgt cgt ttc cag aag ctc gtc gag 1314  
Gly Thr Arg Asp Cys Ser Leu Gln Arg Arg Phe Gln Lys Leu Val Glu  
235 240 245

gag gca ccg gca ccg ttc ctc acc gat gag cag cgt gac cgc atc cac 1362  
Glu Ala Pro Ala Pro Phe Leu Thr Asp Glu Gln Arg Asp Arg Ile His  
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tcc tcc gcc aag gct atc tgc cgc gag gcc ggt tac tac ggt gcc ggc 1410  
Ser Ser Ala Lys Ala Ile Cys Arg Glu Ala Gly Tyr Tyr Gly Ala Gly  
265 270 275

acc gtg gag tac ctg gtc ggt tcc gac gga ctg atc tcc ttc ctg gag 1458  
Thr Val Glu Tyr Leu Val Gly Ser Asp Gly Leu Ile Ser Phe Leu Glu  
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gtc aac acc cgc ctg cag gtg gag cac ccc gtc acc gag gag acc acc 1506  
Val Asn Thr Arg Leu Gln Val Glu His Pro Val Thr Glu Glu Thr Thr  
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Gly Ile Asp Leu Val Arg Glu Met Phe Arg Ile Ala Glu Gly Ala Glu  
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ctc tcc atc aag gag gac ccg acc cca cgc ggc cac gcc ttc gag ttc 1602  
Leu Ser Ile Lys Glu Asp Pro Thr Pro Arg Gly His Ala Phe Glu Phe  
330 335 340

cgc atc aac ggc gag gac gca ggc tcc aac ttc atg ccc gca ccg ggc 1650  
Arg Ile Asn Gly Glu Asp Ala Gly Ser Asn Phe Met Pro Ala Pro Gly  
345 350 355

aag atc acc cgc tac cgt gag ccc gcc ggc ccg ggt gtc cgc atg gac 1698  
Lys Ile Thr Arg Tyr Arg Glu Pro Ala Gly Pro Gly Val Arg Met Asp  
360 365 370

tcc ggc gtt gtc gag ggt tcc gag atc tcc ggc cag ttc gac tcc atg 1746  
Ser Gly Val Val Glu Gly Ser Glu Ile Ser Gly Gln Phe Asp Ser Met  
375 380 385 390

ctg gcc aag ctg atc gtc tgg ggc cag acc cgt gag cag gcc ctg gag 1794  
Leu Ala Lys Leu Ile Val Trp Gly Gln Thr Arg Glu Gln Ala Leu Glu  
395 400 405

cgt tcc cgt cgt gcg ctc ggc gag tac atc gtc gag ggc atg ccg acc 1842  
Arg Ser Arg Arg Ala Leu Gly Glu Tyr Ile Val Glu Gly Met Pro Thr  
410 415 420

gtc atc ccg ttc cac tcc cac atc gtc tcc aac ccg gca ttc gtc ggt 1890  
Val Ile Pro Phe His Ser His Ile Val Ser Asn Pro Ala Phe Val Gly  
425 430 435

gac ggc gag ggc ttc gag gtc tac acc aag tgg atc gag gag gtc tgg 1938  
Asp Gly Glu Gly Phe Glu Val Tyr Thr Lys Trp Ile Glu Glu Val Trp  
440 445 450

gac aac ccg atc gag ccg ttc gtc gat gca gcc gac ctc gac gag 1986

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Asp Asn Pro Ile Glu Pro Phe Val Asp Ala Ala Asp Leu Asp Asp Glu  
 455 460 465 470  
 gag aag acc ccg tcg cag aag gtc atc gtc gag atc gac ggc cgc cgc 2034  
 Glu Lys Thr Pro Ser Gln Lys Val Ile Val Glu Ile Asp Gly Arg Arg  
 475 480 485  
 gtc gag gtg gct ctc ccg ggc gac ctc gct ctc ggc ggt ggc gca ggt 2082  
 Val Glu Val Ala Leu Pro Gly Asp Leu Ala Leu Gly Gly Gly Ala Gly  
 490 495 500  
 gcc gcc aag aag aag ccg aag aag cgt cgc gca ggt ggc gcc aag gcc 2130  
 Ala Ala Lys Lys Lys Pro Lys Lys Arg Arg Ala Gly Gly Ala Lys Ala  
 505 510 515  
 ggt gtc tcc ggt gac tcc gtc gca gcc ccg atg cag ggc acc gtc atc 2178  
 Gly Val Ser Gly Asp Ser Val Ala Ala Pro Met Gln Gly Thr Val Ile  
 520 525 530  
 aag gtc aac gtt gag gac ggc gcc gag gtc tcc gag ggt gac acc gtc 2226  
 Lys Val Asn Val Glu Asp Gly Ala Glu Val Ser Glu Gly Asp Thr Val  
 535 540 545 550  
 gtg gtt ctc gag gcc atg aag atg gag aac ccg gtc aag gcc cac aag 2274  
 Val Val Leu Glu Ala Met Lys Met Glu Asn Pro Val Lys Ala His Lys  
 555 560 565  
 tcc ggt acc gtc tcc ggt ctg acc atc gcc gcg ggt gag ggc gtg acc 2322  
 Ser Gly Thr Val Ser Gly Leu Thr Ile Ala Ala Gly Glu Gly Val Thr  
 570 575 580  
 aag ggt cag gtt ctc ctg gag atc aag taatcccttc agggaacaga 2369  
 Lys Gly Gln Val Leu Leu Glu Ile Lys  
 585 590  
 cagccctgtt ct 2381

&lt;210&gt; 4

&lt;211&gt; 591

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 4

Val Ser Val Glu Thr Arg Lys Ile Thr Lys Val Leu Val Ala Asn Arg  
 1 5 10 15  
 Gly Glu Ile Ala Ile Arg Val Phe Arg Ala Ala Arg Asp Glu Gly Ile  
 20 25 30  
 Ala Ser Val Ala Val Tyr Ala Glu Pro Asp Ala Asp Ala Pro Phe Val  
 35 40 45  
 Glu Tyr Ala Asp Glu Ala Phe Ala Leu Gly Gly Gln Thr Ser Ala Glu  
 50 55 60  
 Ser Tyr Leu Val Ile Asp Lys Ile Ile Asp Ala Ala Arg Lys Ser Gly

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65		70		75		80
Ala Asp Ala Val His Pro Gly Tyr Gly Phe Leu Ala Glu Asn Ala Asp						
	85			90		95
Phe Ala Glu Ala Val Ile Asn Glu Gly Leu Ile Trp Ile Gly Pro Ser						
	100			105		110
Pro Glu Ser Ile Arg Ser Leu Gly Asp Lys Val Thr Ala Arg His Ile						
	115			120		125
Ala Asn Asn Ala Asn Ala Pro Met Ala Pro Gly Thr Lys Glu Pro Val						
	130			135		140
Lys Asp Ala Ala Glu Val Val Ala Phe Ala Glu Glu Phe Gly Leu Pro						
145		150		155		160
Ile Ala Ile Lys Ala Ala Phe Gly Gly Gly Gly Arg Gly Met Lys Val						
	165			170		175
Ala Tyr Glu Met Asp Glu Val Ala Asp Leu Phe Glu Ser Ala Thr Arg						
	180			185		190
Glu Ala Thr Ala Ala Phe Gly Arg Gly Glu Cys Phe Val Glu Arg Tyr						
	195			200		205
Leu Asp Lys Ala Arg His Val Glu Ala Gln Val Ile Ala Asp Lys His						
	210			215		220
Gly Asn Val Val Val Ala Gly Thr Arg Asp Cys Ser Leu Gln Arg Arg						
225		230		235		240
Phe Gln Lys Leu Val Glu Glu Ala Pro Ala Pro Phe Leu Thr Asp Glu						
	245			250		255
Gln Arg Asp Arg Ile His Ser Ser Ala Lys Ala Ile Cys Arg Glu Ala						
	260			265		270
Gly Tyr Tyr Gly Ala Gly Thr Val Glu Tyr Leu Val Gly Ser Asp Gly						
	275			280		285
Leu Ile Ser Phe Leu Glu Val Asn Thr Arg Leu Gln Val Glu His Pro						
	290			295		300
Val Thr Glu Glu Thr Thr Gly Ile Asp Leu Val Arg Glu Met Phe Arg						
305		310		315		320
Ile Ala Glu Gly Ala Glu Leu Ser Ile Lys Glu Asp Pro Thr Pro Arg						
	325			330		335
Gly His Ala Phe Glu Phe Arg Ile Asn Gly Glu Asp Ala Gly Ser Asn						
	340			345		350
Phe Met Pro Ala Pro Gly Lys Ile Thr Arg Tyr Arg Glu Pro Ala Gly						
	355			360		365
Pro Gly Val Arg Met Asp Ser Gly Val Val Glu Gly Ser Glu Ile Ser						
	370			375		380
Gly Gln Phe Asp Ser Met Leu Ala Lys Leu Ile Val Trp Gly Gln Thr						
385		390		395		400
Arg Glu Gln Ala Leu Glu Arg Ser Arg Arg Ala Leu Gly Glu Tyr Ile						
	405			410		415

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Val Glu Gly Met Pro Thr Val Ile Pro Phe His Ser His Ile Val Ser  
                     420                    425                    430  
 Asn Pro Ala Phe Val Gly Asp Gly Glu Gly Phe Glu Val Tyr Thr Lys  
                     435                    440                    445  
 Trp Ile Glu Glu Val Trp Asp Asn Pro Ile Glu Pro Phe Val Asp Ala  
                     450                    455                    460  
 Ala Asp Leu Asp Asp Glu Glu Lys Thr Pro Ser Gln Lys Val Ile Val  
 465                    470                    475                    480  
 Glu Ile Asp Gly Arg Arg Val Glu Val Ala Leu Pro Gly Asp Leu Ala  
                     485                    490                    495  
 Leu Gly Gly Gly Ala Gly Ala Ala Lys Lys Lys Pro Lys Lys Arg Arg  
                     500                    505                    510  
 Ala Gly Gly Ala Lys Ala Gly Val Ser Gly Asp Ser Val Ala Ala Pro  
                     515                    520                    525  
 Met Gln Gly Thr Val Ile Lys Val Asn Val Glu Asp Gly Ala Glu Val  
                     530                    535                    540  
 Ser Glu Gly Asp Thr Val Val Val Leu Glu Ala Met Lys Met Glu Asn  
 545                    550                    555                    560  
 Pro Val Lys Ala His Lys Ser Gly Thr Val Ser Gly Leu Thr Ile Ala  
                     565                    570                    575  
 Ala Gly Glu Gly Val Thr Lys Gly Gln Val Leu Leu Glu Ile Lys  
                     580                    585                    590

<210> 5  
 <211> 2128  
 <212> DNA  
 <213> Corynebacterium thermoaminogenes

<220>  
 <221> CDS  
 <222> (339).. (1967)

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 atccacctgt ggaacagtca gcggcgcggc catggagggc agcgacaggt gacgtccgag 180  
 caccgggttc cccaccgigg acacggcatt gatccgacac ggtggggata gtltcatgct 240  
 gaaaaactat cgtgtgtcag ggaggatccg gaatgtgacc tatltcatgg agaaatgatt 300  
 gtggacgata cccccgggta cggctacctt tccaaaac atg acc att tcc tca cct 356

Met Thr Ile Ser Ser Pro

1

5

ttg att gac gtc gct aac ctg cca gac atc aac acc acc gcc ggc aag 404  
 Leu Ile Asp Val Ala Asn Leu Pro Asp Ile Asn Thr Thr Ala Gly Lys



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gcc acc gat gag gag gcc ctc gac tgg gtc cag gac ctc atc tcc ttc	1124
Ala Thr Asp Glu Glu Ala Leu Asp Trp Val Gln Asp Leu Ile Ser Phe	
250 255 260	
ctg ccc tcc aac aat cgc tcc tac gcc ccg gtg gag gag ttc gac gag	1172
Leu Pro Ser Asn Asn Arg Ser Tyr Ala Pro Val Glu Glu Phe Asp Glu	
265 270 275	
gag gac ggt ggc atc gcc gag aac atc acc gcc gat gac ctg aag ctg	1220
Glu Asp Gly Gly Ile Ala Glu Asn Ile Thr Ala Asp Asp Leu Lys Leu	
280 285 290	
gat gag atc atc ccg gat tcc gcc acc gtg ccc tat gat gtc cgc gac	1268
Asp Glu Ile Ile Pro Asp Ser Ala Thr Val Pro Tyr Asp Val Arg Asp	
295 300 305 310	
gtc atc cag tgc ctg acc gac gac ggt gag tac ctg gag atc cag gcc	1316
Val Ile Gln Cys Leu Thr Asp Asp Gly Glu Tyr Leu Glu Ile Gln Ala	
315 320 325	
gac cga gcc gag aat gtc gtc atc gcc ttc ggc cgc atc gag ggc cag	1364
Asp Arg Ala Glu Asn Val Val Ile Ala Phe Gly Arg Ile Glu Gly Gln	
330 335 340	
tcc gtc ggt ttc gtc gcc aac cag ccg acc cag ttc gcc ggc tgc ctg	1412
Ser Val Gly Phe Val Ala Asn Gln Pro Thr Gln Phe Ala Gly Cys Leu	
345 350 355	
gac atc gac tcc tcc gag aag gca gcc cgc ttc gtc cgc acc tgc gat	1460
Asp Ile Asp Ser Ser Glu Lys Ala Ala Arg Phe Val Arg Thr Cys Asp	
360 365 370	
gcc ttc aac atc ccg atc gtc atg ctt gtc gac gtc ccc ggc ttc ctc	1508
Ala Phe Asn Ile Pro Ile Val Met Leu Val Asp Val Pro Gly Phe Leu	
375 380 385 390	
ccc ggt gcc ggc cag gag tac ggc ggc atc ctg cgt cgt ggc gcc aaa	1556
Pro Gly Ala Gly Gln Glu Tyr Gly Gly Ile Leu Arg Arg Gly Ala Lys	
395 400 405	
ctg ctc tac gcc tac ggt gag gcc acc gtc ccg aag atc acc gtg acc	1604
Leu Leu Tyr Ala Tyr Gly Glu Ala Thr Val Pro Lys Ile Thr Val Thr	
410 415 420	
atg cgc aag gcc tac ggc ggt gcg tac tgt gtc atg gga tcc aag ggt	1652
Met Arg Lys Ala Tyr Gly Gly Ala Tyr Cys Val Met Gly Ser Lys Gly	
425 430 435	
ctg ggc gca gac atc aac ctg gcc tgg ccg acc gcg cag atc gcc gtc	1700
Leu Gly Ala Asp Ile Asn Leu Ala Trp Pro Thr Ala Gln Ile Ala Val	
440 445 450	
atg ggt gcc gcc ggc gcg gtc cag ttc atc tac cgc aag gag ctc atg	1748
Met Gly Ala Ala Gly Ala Val Gln Phe Ile Tyr Arg Lys Glu Leu Met	
455 460 465 470	
gcc gct gat gcc aag ggc ctg gac acc gtc gcc ctg gcc cag tcc ttc	1796

Met	Thr	Ile	Ser	Ser	Pro	Leu	Ile	Asp	Val	Ala	Asn	Leu	Pro	Asp	Ile
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			20					25					30		
Ala	His	Phe	Pro	Met	Gly	Glu	Lys	Ala	Val	Glu	Lys	Val	His	Ala	Ala
		35					40						45		
Asn	Arg	Leu	Thr	Ala	Arg	Glu	Arg	Leu	Asp	Tyr	Leu	Leu	Asp	Glu	Gly
	50					55					60				
Ser	Phe	Ile	Glu	Thr	Asp	Gln	Leu	Ala	Arg	His	Arg	Thr	Thr	Ala	Phe
65					70					75					80
Gly	Leu	Gly	Asn	Lys	Arg	Pro	Ala	Thr	Asp	Gly	Ile	Val	Thr	Gly	Trp
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Gly	Thr	Ile	Asp	Gly	Arg	Glu	Val	Cys	Ile	Phe	Ser	Gln	Asp	Gly	Thr
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Val	Phe	Gly	Gly	Ala	Leu	Gly	Glu	Val	Tyr	Gly	Glu	Lys	Met	Ile	Lys
		115					120					125			
Ile	Met	Glu	Leu	Ala	Ile	Asp	Thr	Gly	Arg	Pro	Leu	Ile	Gly	Leu	Tyr
	130					135					140				
Glu	Gly	Ala	Gly	Ala	Arg	Ile	Gln	Asp	Gly	Ala	Val	Ser	Leu	Asp	Phe



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145		150		155		160									
Ile	Ser	Gln	Thr	Phe	Tyr	Gln	Asn	Ile	Gln	Ala	Ser	Gly	Val	Ile	Pro
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Gln	Ile	Ser	Val	Ile	Met	Gly	Ala	Cys	Ala	Gly	Gly	Asn	Ala	Tyr	Gly
		180				185							190		
Pro	Ala	Leu	Thr	Asp	Phe	Val	Val	Met	Val	Asp	Lys	Thr	Ser	Lys	Met
		195				200						205			
Phe	Val	Thr	Gly	Pro	Asp	Val	Ile	Lys	Thr	Val	Thr	Gly	Glu	Glu	Ile
		210				215					220				
Thr	Gln	Glu	Glu	Leu	Gly	Gly	Ala	Thr	Thr	His	Met	Val	Thr	Ala	Gly
225					230					235				240	
Asn	Ser	His	Tyr	Thr	Val	Ala	Thr	Asp	Glu	Glu	Ala	Leu	Asp	Trp	Val
			245					250					255		
Gln	Asp	Leu	Ile	Ser	Phe	Leu	Pro	Ser	Asn	Asn	Arg	Ser	Tyr	Ala	Pro
		260				265							270		
Val	Glu	Glu	Phe	Asp	Glu	Glu	Asp	Gly	Gly	Ile	Ala	Glu	Asn	Ile	Thr
		275				280					285				
Ala	Asp	Asp	Leu	Lys	Leu	Asp	Glu	Ile	Ile	Pro	Asp	Ser	Ala	Thr	Val
		290				295					300				
Pro	Tyr	Asp	Val	Arg	Asp	Val	Ile	Gln	Cys	Leu	Thr	Asp	Asp	Gly	Glu
305				310					315					320	
Tyr	Leu	Glu	Ile	Gln	Ala	Asp	Arg	Ala	Glu	Asn	Val	Val	Ile	Ala	Phe
			325					330					335		
Gly	Arg	Ile	Glu	Gly	Gln	Ser	Val	Gly	Phe	Val	Ala	Asn	Gln	Pro	Thr
		340						345					350		
Gln	Phe	Ala	Gly	Cys	Leu	Asp	Ile	Asp	Ser	Ser	Glu	Lys	Ala	Ala	Arg
		355				360					365				
Phe	Val	Arg	Thr	Cys	Asp	Ala	Phe	Asn	Ile	Pro	Ile	Val	Met	Leu	Val
		370				375					380				
Asp	Val	Pro	Gly	Phe	Leu	Pro	Gly	Ala	Gly	Gln	Glu	Tyr	Gly	Gly	Ile
385				390						395				400	
Leu	Arg	Arg	Gly	Ala	Lys	Leu	Leu	Tyr	Ala	Tyr	Gly	Glu	Ala	Thr	Val
			405					410					415		
Pro	Lys	Ile	Thr	Val	Thr	Met	Arg	Lys	Ala	Tyr	Gly	Gly	Ala	Tyr	Cys
		420						425					430		
Val	Met	Gly	Ser	Lys	Gly	Leu	Gly	Ala	Asp	Ile	Asn	Leu	Ala	Trp	Pro
		435				440					445				
Thr	Ala	Gln	Ile	Ala	Val	Met	Gly	Ala	Ala	Gly	Ala	Val	Gln	Phe	Ile
		450				455					460				
Tyr	Arg	Lys	Glu	Leu	Met	Ala	Ala	Asp	Ala	Lys	Gly	Leu	Asp	Thr	Val
465				470						475				480	
Ala	Leu	Ala	Gln	Ser	Phe	Glu	Arg	Glu	Tyr	Glu	Asp	His	Met	Leu	Asn
			485					490					495		

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Pro Tyr Leu Ala Ala Glu Arg Gly Leu Ile Asp Ala Val Ile Leu Pro  
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Ser Glu Thr Arg Gly Gln Ile Ala Arg Asn Leu Arg Leu Leu Lys His  
515 520 525

Lys Asn Val Ser Arg Pro Ala Arg Lys His Gly Asn Met Pro Leu  
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<210>	7
<211>	2076
<212>	DNA
<213>	Corynebacterium thermoaminogenes

$\langle 220 \rangle$   
 $\langle 221 \rangle$  CDS  
 $\langle 222 \rangle$  (412) .. (2022)

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acggggggga	ggagggtaca	taggccatac	gtcgcacttt	tgatgaagtg	tgggcagatc	180									
gaccgggcaa	atcigggaaa	taagggggcc	ggtgaactag	cattccccct	agcgaagggt	240									
gagcatcgcg	gaccccgcg	gttcccaacc	ggtcglaaat	tcatgtgccg	ccacagtcct	300									
ctcaccaggg	galtcggaacc	agcccagcc	gattccggcg	tgacggacct	caccgigaac	360									
aagtcctcgc	attactcaca	gaattcacac	caggattttg	actaagaaac	c atg act	417									
					Met Thr										
					1										
gca gca acg aca gca cct gat ctg acc acc acc gcc ggc aaa ctc gcg						465									
Ala Ala Thr Thr Ala Pro Asp Leu Thr Thr Thr Ala Gly Lys Leu Ala															
	5														
gat ctc cgc gcc cgc ctt tcc gag acc cag gcc ccc atg ggt cag gcc						513									
Asp Leu Arg Ala Arg Leu Ser Glu Thr Gln Ala Pro Met Gly Gln Ala															
	20														
tcc gtg gag aag gtg cac gag gca ggg aag aag acc gca cgc gag cgc						561									
Ser Val Glu Lys Val His Glu Ala Gly Lys Lys Thr Ala Arg Glu Arg															
	35														
atc gag tac ctg ctc gat gag ggc tcc ttc gtt gag gtc gat gcc ctc						609									
Ile Glu Tyr Leu Leu Asp Glu Gly Ser Phe Val Glu Val Asp Ala Leu															
gcc cgc cac cgt tcc aag aac ttc ggc ctg gac tcc aag cgc ccg gtc						657									
Ala Arg His Arg Ser Lys Asn Phe Gly Leu Asp Ser Lys Arg Pro Val															
acc gac ggt gtg gtc acc ggt tac ggc acc atc gac gga cgc aag gtc						705									
Thr Asp Gly Val Val Thr Gly Tyr Gly Thr Ile Asp Gly Arg Lys Val															

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85	90	95	
tgc gtc ttc tcc cag gac ggc gct atc ttc ggc ggt gcc ctc ggt gag			753
Cys Val Phe Ser Gln Asp Gly Ala Ile Phe Gly Gly Ala Leu Gly Glu			
100	105	110	
gtc tac ggc gag aag atc gtc aag atc atg gac ctg gcc atc aag acc			801
Val Tyr Gly Glu Lys Ile Val Lys Ile Met Asp Leu Ala Ile Lys Thr			
115	120	125	130
ggt gtc ccc ctc atc ggc atc aac gag ggc gcc ggc gcc cgc atc cag			849
Gly Val Pro Leu Ile Gly Ile Asn Glu Gly Ala Gly Ala Arg Ile Gln			
135	140	145	
gaa ggc gtt gtc tcc ctg ggc ctg tac tcc cag atc ttc tac cgc aac			897
Glu Gly Val Val Ser Leu Gly Leu Tyr Ser Gln Ile Phe Tyr Arg Asn			
150	155	160	
acc cag gca tcc ggt gtc atc cca cag atc tcc ctc atc atg ggt gcc			945
Thr Gln Ala Ser Gly Val Ile Pro Gln Ile Ser Leu Ile Met Gly Ala			
165	170	175	
tgc gcc ggt ggc cat gtg tac tcc ccc gcc ctg acc gac ttc atc atc			993
Cys Ala Gly Gly His Val Tyr Ser Pro Ala Leu Thr Asp Phe Ile Ile			
180	185	190	
atg gtg gac aag acc tcc aag atg ttc atc acc ggc ccc gac gtg atc			1041
Met Val Asp Lys Thr Ser Lys Met Phe Ile Thr Gly Pro Asp Val Ile			
195	200	205	210
aag acc gtc acc ggc gag gag gtc acc cag gag gaa ctg ggt ggt gcc			1089
Lys Thr Val Thr Gly Glu Glu Val Thr Gln Glu Glu Leu Gly Gly Ala			
215	220	225	
tac acc cac atg gcc cag tcc ggc acc tgc cac tac acc gca gcc gat			1137
Tyr Thr His Met Ala Gln Ser Gly Thr Ser His Tyr Thr Ala Ala Asp			
230	235	240	
gac tcc gat gcc ctc gac tgg gtc cgt gag ctg gtc agc tac ctg ccg			1185
Asp Ser Asp Ala Leu Asp Trp Val Arg Glu Leu Val Ser Tyr Leu Pro			
245	250	255	
tcc aac aac cgt gcg gag acc cca cgc cag gac gcc gac atc atg gtg			1233
Ser Asn Asn Arg Ala Glu Thr Pro Arg Gln Asp Ala Asp Ile Met Val			
260	265	270	
ggc tcc atc aag gag aac atc acc gag acc gac ctc gaa ctc gac acc			1281
Gly Ser Ile Lys Glu Asn Ile Thr Glu Thr Asp Leu Glu Leu Asp Thr			
275	280	285	290
ctg atc ccg gat tcc ccg aac cag ccg tac gac atg aag gac gtc atc			1329
Leu Ile Pro Asp Ser Pro Asn Gln Pro Tyr Asp Met Lys Asp Val Ile			
295	300	305	
acc cgc atc gtc gat gat gcc gag ttc ttc gag atc cag gag ggt tac			1377
Thr Arg Ile Val Asp Asp Ala Glu Phe Phe Glu Ile Gln Glu Gly Tyr			
310	315	320	

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gcc gag aac atc atc tgc ggt ttc gcc cgc gtc gag ggt cgt gcc gtg	1425
Ala Glu Asn Ile Ile Cys Gly Phe Ala Arg Val Glu Gly Arg Ala Val	
325 330 335	
ggt atc gtg gcc aac cag ccg atg cag ttc gcc ggc tgc ctg gac atc	1473
Gly Ile Val Ala Asn Gln Pro Met Gln Phe Ala Gly Cys Leu Asp Ile	
340 345 350	
aag gca tcc gag aag gcc gcc cgc ttc atc cgc acc tgt gac gcc ttc	1521
Lys Ala Ser Glu Lys Ala Ala Arg Phe Ile Arg Thr Cys Asp Ala Phe	
355 360 365 370	
aac atc ccg atc atc gag ctt gtc gac gtc cca ggc ttc ctc ccg ggc	1569
Asn Ile Pro Ile Ile Glu Leu Val Asp Val Pro Gly Phe Leu Pro Gly	
375 380 385	
acc aac cag gag ttc gac ggc atc atc cgt cgc ggc gcg aag ctg ctc	1617
Thr Asn Gln Glu Phe Asp Gly Ile Ile Arg Arg Gly Ala Lys Leu Leu	
390 395 400	
tac gcc tac gcc gag gcc acc gtc ggc aag atc acc gtg atc acc cgc	1665
Tyr Ala Tyr Ala Glu Ala Thr Val Gly Lys Ile Thr Val Ile Thr Arg	
405 410 415	
aag tcc tac ggc ggt gcc tac tgc gtg atg ggc tcc aag gac atg ggt	1713
Lys Ser Tyr Gly Gly Ala Tyr Cys Val Met Gly Ser Lys Asp Met Gly	
420 425 430	
gcg gac ctc gtc ttc gca tgg ccc acc gcg cag atc gcc gtc atg ggt	1761
Ala Asp Leu Val Phe Ala Trp Pro Thr Ala Gln Ile Ala Val Met Gly	
435 440 445 450	
gcc tcc ggt gcc gtc ggc ttc atc tac cgc aag gag ctc aag cag gct	1809
Ala Ser Gly Ala Val Gly Phe Ile Tyr Arg Lys Glu Leu Lys Gln Ala	
455 460 465	
gca gcg gcc ggc gag gat gtc acc gcg ctg atg aag aag tac gag cag	1857
Ala Ala Ala Gly Glu Asp Val Thr Ala Leu Met Lys Lys Tyr Glu Gln	
470 475 480	
gag tac gag gag acc ctg gtc aac ccg tac atg gct gca gag cgt ggc	1905
Glu Tyr Glu Glu Thr Leu Val Asn Pro Tyr Met Ala Ala Glu Arg Gly	
485 490 495	
tac gtc gac gcc gtc atc cca cca tcc gag acc cgt ggt cag atc atc	1953
Tyr Val Asp Ala Val Ile Pro Pro Ser Glu Thr Arg Gly Gln Ile Ile	
500 505 510	
gag ggt ctg cgt ctg ctc gac cgc aag gtg gtc aac gtc ccg gcc aag	2001
Glu Gly Leu Arg Leu Leu Asp Arg Lys Val Val Asn Val Pro Ala Lys	
515 520 525 530	
aag cac ggt aac atc ccg ctg taaaccgtct tccctccgg caccacgccg	2052
Lys His Gly Asn Ile Pro Leu	
535	
gagaaggctt tglccgcagc tgtc	2076

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&lt;210&gt; 8

&lt;211&gt; 537

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 8

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Met Thr Ala Ala Thr Thr Ala Pro Asp Leu Thr Thr Thr Ala Gly Lys
 1           5           10           15
Leu Ala Asp Leu Arg Ala Arg Leu Ser Glu Thr Gln Ala Pro Met Gly
      20           25           30
Gln Ala Ser Val Glu Lys Val His Glu Ala Gly Lys Lys Thr Ala Arg
      35           40           45
Glu Arg Ile Glu Tyr Leu Leu Asp Glu Gly Ser Phe Val Glu Val Asp
      50           55           60
Ala Leu Ala Arg His Arg Ser Lys Asn Phe Gly Leu Asp Ser Lys Arg
      65           70           75           80
Pro Val Thr Asp Gly Val Val Thr Gly Tyr Gly Thr Ile Asp Gly Arg
      85           90           95
Lys Val Cys Val Phe Ser Gln Asp Gly Ala Ile Phe Gly Gly Ala Leu
      100          105          110
Gly Glu Val Tyr Gly Glu Lys Ile Val Lys Ile Met Asp Leu Ala Ile
      115          120          125
Lys Thr Gly Val Pro Leu Ile Gly Ile Asn Glu Gly Ala Gly Ala Arg
      130          135          140
Ile Gln Glu Gly Val Val Ser Leu Gly Leu Tyr Ser Gln Ile Phe Tyr
      145          150          155          160
Arg Asn Thr Gln Ala Ser Gly Val Ile Pro Gln Ile Ser Leu Ile Met
      165          170          175
Gly Ala Cys Ala Gly Gly His Val Tyr Ser Pro Ala Leu Thr Asp Phe
      180          185          190
Ile Ile Met Val Asp Lys Thr Ser Lys Met Phe Ile Thr Gly Pro Asp
      195          200          205
Val Ile Lys Thr Val Thr Gly Glu Glu Val Thr Gln Glu Glu Leu Gly
      210          215          220
Gly Ala Tyr Thr His Met Ala Gln Ser Gly Thr Ser His Tyr Thr Ala
      225          230          235          240
Ala Asp Asp Ser Asp Ala Leu Asp Trp Val Arg Glu Leu Val Ser Tyr
      245          250          255
Leu Pro Ser Asn Asn Arg Ala Glu Thr Pro Arg Gln Asp Ala Asp Ile
      260          265          270
Met Val Gly Ser Ile Lys Glu Asn Ile Thr Glu Thr Asp Leu Glu Leu
      275          280          285

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Asp Thr Leu Ile Pro Asp Ser Pro Asn Gln Pro Tyr Asp Met Lys Asp  
 290 295 300  
 Val Ile Thr Arg Ile Val Asp Asp Ala Glu Phe Phe Glu Ile Gln Glu  
 305 310 315 320  
 Gly Tyr Ala Glu Asn Ile Ile Cys Gly Phe Ala Arg Val Glu Gly Arg  
 325 330 335  
 Ala Val Gly Ile Val Ala Asn Gln Pro Met Gln Phe Ala Gly Cys Leu  
 340 345 350  
 Asp Ile Lys Ala Ser Glu Lys Ala Ala Arg Phe Ile Arg Thr Cys Asp  
 355 360 365  
 Ala Phe Asn Ile Pro Ile Ile Glu Leu Val Asp Val Pro Gly Phe Leu  
 370 375 380  
 Pro Gly Thr Asn Gln Glu Phe Asp Gly Ile Ile Arg Arg Gly Ala Lys  
 385 390 395 400  
 Leu Leu Tyr Ala Tyr Ala Glu Ala Thr Val Gly Lys Ile Thr Val Ile  
 405 410 415  
 Thr Arg Lys Ser Tyr Gly Gly Ala Tyr Cys Val Met Gly Ser Lys Asp  
 420 425 430  
 Met Gly Ala Asp Leu Val Phe Ala Trp Pro Thr Ala Gln Ile Ala Val  
 435 440 445  
 Met Gly Ala Ser Gly Ala Val Gly Phe Ile Tyr Arg Lys Glu Leu Lys  
 450 455 460  
 Gln Ala Ala Ala Ala Gly Glu Asp Val Thr Ala Leu Met Lys Lys Tyr  
 465 470 475 480  
 Glu Gln Glu Tyr Glu Glu Thr Leu Val Asn Pro Tyr Met Ala Ala Glu  
 485 490 495  
 Arg Gly Tyr Val Asp Ala Val Ile Pro Pro Ser Glu Thr Arg Gly Gln  
 500 505 510  
 Ile Ile Glu Gly Leu Arg Leu Leu Asp Arg Lys Val Val Asn Val Pro  
 515 520 525  
 Ala Lys Lys His Gly Asn Ile Pro Leu  
 530 535

&lt;210&gt; 9

&lt;211&gt; 1643

&lt;212&gt; DNA

<213> *Corynebacterium thermoaminogenes*

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (326)..(1363)

&lt;400&gt; 9

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agcgcgccgg cagccaccag tgggatcgig cccagcggac ggaigccgga ttcacggcgg 60  
 tcagccaccc gccgatgaga cctgcagcga caacggtggc ggtgcigacc tggtcagcgt 120  
 cttigagttt cataatcaig tcagacagtc taaccactct ctccgacgcg tccgaacaig 180  
 ctgggggtggc ggacaccaig tccgttcggg cggtgccccg acgggggaaa atcgcaggca 240  
 gatgltccg atgigggata aaccaccgg ttcgggcgtg tcttcgggat caatggcaca 300  
 gcatiaaccg tgtgggggggt ttaat atg gga gcc atg cga att gcc act ctc 352  
 Met Gly Ala Met Arg Ile Ala Thr Leu  
 1 5  
 acg tca ggc ggc gac tgc ccc gga ctc aat gct gtc atc agg gga atc 400  
 Thr Ser Gly Gly Asp Cys Pro Gly Leu Asn Ala Val Ile Arg Gly Ile  
 10 15 20 25  
 gtc cgt acc gca agt aat gaa ttc ggt tcc acc gtc gtg ggt tat cag 448  
 Val Arg Thr Ala Ser Asn Glu Phe Gly Ser Thr Val Val Gly Tyr Gln  
 30 35 40  
 gac ggc tgg gag ggc ctg ctg gcg gac cga cgt gtt cag ctc tat gac 496  
 Asp Gly Trp Glu Gly Leu Leu Ala Asp Arg Arg Val Gln Leu Tyr Asp  
 45 50 55  
 gat gag gac atc gac cgc atc ctg ctc cgc ggt gga aca atc ctg ggc 544  
 Asp Glu Asp Ile Asp Arg Ile Leu Leu Arg Gly Gly Thr Ile Leu Gly  
 60 65 70  
 acc ggt cgt ctc cac ccc gac aag ttc aga gcc gga atc gac cag gtc 592  
 Thr Gly Arg Leu His Pro Asp Lys Phe Arg Ala Gly Ile Asp Gln Val  
 75 80 85  
 aag gcg aat ctc gcc gat gcg gga att gac gca ctc atc ccg atc ggt 640  
 Lys Ala Asn Leu Ala Asp Ala Gly Ile Asp Ala Leu Ile Pro Ile Gly  
 90 95 100 105  
 ggc gag ggc acc ctc aag gga gcg aag tgg ctc gcc gac aac ggc atc 688  
 Gly Glu Gly Thr Leu Lys Gly Ala Lys Trp Leu Ala Asp Asn Gly Ile  
 110 115 120  
 ccc gtg gtc ggt gtc ccg aaa acc atc gac aat gat gtc aac ggc acg 736  
 Pro Val Val Gly Val Pro Lys Thr Ile Asp Asn Asp Val Asn Gly Thr  
 125 130 135  
 gat ttc acc ttc ggt ttc gat tcc gcg gtc tct gtg gcc acc gac gcc 784  
 Asp Phe Thr Phe Gly Phe Asp Ser Ala Val Ser Val Ala Thr Asp Ala  
 140 145 150  
 atc gac cgg ctg cac acc acg gcg gaa tcc cac aac cgt gtg atg atc 832  
 Ile Asp Arg Leu His Thr Thr Ala Glu Ser His Asn Arg Val Met Ile  
 155 160 165  
 gtc gag gtc atg ggc cgc cac gtc ggt tgg atc gca ctg cat gcc ggc 880  
 Val Glu Val Met Gly Arg His Val Gly Trp Ile Ala Leu His Ala Gly  
 170 175 180 185  
 atg gcc ggt gga gcc cac tac acc gtc atc ccc gag gtg ccc ttc gac 928  
 Met Ala Gly Gly Ala His Tyr Thr Val Ile Pro Glu Val Pro Phe Asp

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190	195	200	
atc tgc gag atc tgc aag cgt atg gaa cgt cgc ttc cag atg ggg gag			976
Ile Ser Glu Ile Cys Lys Arg Met Glu Arg Arg Phe Gln Met Gly Glu			
205	210	215	
aag tac ggc atc atc gtc gtc gcg gag ggt gcc ctg ccc aag gag gga			1024
Lys Tyr Gly Ile Ile Val Val Ala Glu Gly Ala Leu Pro Lys Glu Gly			
220	225	230	
acc atg gag ctg cgt gag ggg gag gtg gat cag ttc ggt cac aag acc			1072
Thr Met Glu Leu Arg Glu Gly Glu Val Asp Gln Phe Gly His Lys Thr			
235	240	245	
ttc acc ggc atc ggc cag cag atc gcc gac gag gtg cac agg cgt ctg			1120
Phe Thr Gly Ile Gly Gln Gln Ile Ala Asp Glu Val His Arg Arg Leu			
250	255	260	265
ggt cat gat gtc cgc acc acg gtc ctg ggc cat atc cag cgt ggt ggc			1168
Gly His Asp Val Arg Thr Thr Val Leu Gly His Ile Gln Arg Gly Gly			
270	275	280	
acc ccc acc gcc ttc gac cgt gtc ctg gcc acc cgg tac ggt gtc cgc			1216
Thr Pro Thr Ala Phe Asp Arg Val Leu Ala Thr Arg Tyr Gly Val Arg			
285	290	295	
gcc gcg cgt gcc tgc cac gag ggt cag ttc aac acc gtg gtc gcg ctc			1264
Ala Ala Arg Ala Cys His Glu Gly Gln Phe Asn Thr Val Val Ala Leu			
300	305	310	
aag ggg gag cgc atc cgg atg atc tcc ttc gat gag gcc glg ggc acc			1312
Lys Gly Glu Arg Ile Arg Met Ile Ser Phe Asp Glu Ala Val Gly Thr			
315	320	325	
ctg aag aag gtg ccg atg gaa cgc tgg gtg acc gcc cag gct atg ttc			1360
Leu Lys Lys Val Pro Met Glu Arg Trp Val Thr Ala Gln Ala Met Phe			
330	335	340	345
ggt tagtcaggcc gcatcccggt ttcgcgcgcc gcggggccgg gttttttcat			1413
Gly			
gccccggaac acatcggtat gaaatcgiga tatgcattac ttgacgggga agtgggggat			1473
ccgtcaccic gcgttgcca actacagccc gcagcgccctg cgggaattct tcgagcaatc			1533
cgccgatccc ccggcccgtc ccgtcgccgt ccaaccgcag tacaatctgc tggccccccg			1593
ggattatgag accggtatcc gcccggtcgt ggacgagttc ggtcccgcg			1643

<210> 10

<211> 346

<212> PRT

<213> Corynebacterium thermoaminogenes

<400> 10

Met Gly Ala Met Arg Ile Ala Thr Leu Thr Ser Gly Gly Asp Cys Pro

1

5

10

15



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Gly Leu Asn Ala Val Ile Arg Gly Ile Val Arg Thr Ala Ser Asn Glu  
                   20                  25                  30  
 Phe Gly Ser Thr Val Val Gly Tyr Gln Asp Gly Trp Glu Gly Leu Leu  
                   35                  40                  45  
 Ala Asp Arg Arg Val Gln Leu Tyr Asp Asp Glu Asp Ile Asp Arg Ile  
                   50                  55                  60  
 Leu Leu Arg Gly Gly Thr Ile Leu Gly Thr Gly Arg Leu His Pro Asp  
                   65                  70                  75                  80  
 Lys Phe Arg Ala Gly Ile Asp Gln Val Lys Ala Asn Leu Ala Asp Ala  
                   85                  90                  95  
 Gly Ile Asp Ala Leu Ile Pro Ile Gly Gly Glu Gly Thr Leu Lys Gly  
                   100                  105                  110  
 Ala Lys Trp Leu Ala Asp Asn Gly Ile Pro Val Val Gly Val Pro Lys  
                   115                  120                  125  
 Thr Ile Asp Asn Asp Val Asn Gly Thr Asp Phe Thr Phe Gly Phe Asp  
                   130                  135                  140  
 Ser Ala Val Ser Val Ala Thr Asp Ala Ile Asp Arg Leu His Thr Thr  
                   145                  150                  155                  160  
 Ala Glu Ser His Asn Arg Val Met Ile Val Glu Val Met Gly Arg His  
                   165                  170                  175  
 Val Gly Trp Ile Ala Leu His Ala Gly Met Ala Gly Gly Ala His Tyr  
                   180                  185                  190  
 Thr Val Ile Pro Glu Val Pro Phe Asp Ile Ser Glu Ile Cys Lys Arg  
                   195                  200                  205  
 Met Glu Arg Arg Phe Gln Met Gly Glu Lys Tyr Gly Ile Ile Val Val  
                   210                  215                  220  
 Ala Glu Gly Ala Leu Pro Lys Glu Gly Thr Met Glu Leu Arg Glu Gly  
                   225                  230                  235                  240  
 Glu Val Asp Gln Phe Gly His Lys Thr Phe Thr Gly Ile Gly Gln Gln  
                   245                  250                  255  
 Ile Ala Asp Glu Val His Arg Arg Leu Gly His Asp Val Arg Thr Thr  
                   260                  265                  270  
 Val Leu Gly His Ile Gln Arg Gly Gly Thr Pro Thr Ala Phe Asp Arg  
                   275                  280                  285  
 Val Leu Ala Thr Arg Tyr Gly Val Arg Ala Ala Arg Ala Cys His Glu  
                   290                  295                  300  
 Gly Gln Phe Asn Thr Val Val Ala Leu Lys Gly Glu Arg Ile Arg Met  
                   305                  310                  315                  320  
 Ile Ser Phe Asp Glu Ala Val Gly Thr Leu Lys Lys Val Pro Met Glu  
                   325                  330                  335  
 Arg Trp Val Thr Ala Gln Ala Met Phe Gly  
                   340                  345

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<210> 11  
 <211> 498  
 <212> DNA  
 <213> *Corynebacterium thermoaminogenes*

<220>  
 <221> CDS  
 <222> (1)..(498)

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 Tyr Tyr Gln His Asp Pro Gly Phe Pro Phe Ala Pro Lys Arg Thr Gly  
 1 5 10 15  
 tgg gct cac acc acc acg ccg ttg acc gga ccg cag cga ttg cag tgg 96  
 Trp Ala His Thr Thr Thr Pro Leu Thr Gly Pro Gln Arg Leu Gln Trp  
 20 25 30  
 acg cac ctg ccc gat gct ctt tac ccg gat gta tcc tat gac ctg gat 144  
 Thr His Leu Pro Asp Ala Leu Tyr Pro Asp Val Ser Tyr Asp Leu Asp  
 35 40 45  
 gga tgc tat tcc ggc gga gcc gta ttt tct gac ggc acg ctt aaa ctt 192  
 Gly Cys Tyr Ser Gly Gly Ala Val Phe Ser Asp Gly Thr Leu Lys Leu  
 50 55 60  
 ttc tac acc ggc aac cga aaa att gac ggc aag cgc cgc gcc acc caa 240  
 Phe Tyr Thr Gly Asn Arg Lys Ile Asp Gly Lys Arg Arg Ala Thr Gln  
 65 70 75 80  
 aac ctc gtc gaa gtc gag gac cca act ggg ctg atg ggc ggc att cat 288  
 Asn Leu Val Glu Val Glu Asp Pro Thr Gly Leu Met Gly Gly Ile His  
 85 90 95  
 cgc cgc tgc cct aaa aat ccg ctt atc gac gga ccc gcc agc ggt ttt 336  
 Arg Arg Ser Pro Lys Asn Pro Leu Ile Asp Gly Pro Ala Ser Gly Phe  
 100 105 110  
 acg ccc cac tac cgc gat ccc atg atc agc cct gat ggg gat ggt tgg 384  
 Thr Pro His Tyr Arg Asp Pro Met Ile Ser Pro Asp Gly Asp Gly Trp  
 115 120 125  
 aag atg gtt ctt ggg gct cag cgc gaa aac ctc acc ggt gca gcg gtt 432  
 Lys Met Val Leu Gly Ala Gln Arg Glu Asn Leu Thr Gly Ala Ala Val  
 130 135 140  
 cta tac cgc tgc gca gat ctt gaa aac tgg gaa ttc tcc ggt gaa atc 480  
 Leu Tyr Arg Ser Ala Asp Leu Glu Asn Trp Glu Phe Ser Gly Glu Ile  
 145 150 155 160  
 acc ttt gac ctc agc gac 498  
 Thr Phe Asp Leu Ser Asp  
 165

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<210> 12  
 <211> 166  
 <212> PRT  
 <213> Corynebacterium thermoaminogenes

<400> 12

Tyr	Tyr	Gln	His	Asp	Pro	Gly	Phe	Pro	Phe	Ala	Pro	Lys	Arg	Thr	Gly
1				5				10						15	
Trp	Ala	His	Thr	Thr	Thr	Pro	Leu	Thr	Gly	Pro	Gln	Arg	Leu	Gln	Trp
			20					25					30		
Thr	His	Leu	Pro	Asp	Ala	Leu	Tyr	Pro	Asp	Val	Ser	Tyr	Asp	Leu	Asp
		35					40					45			
Gly	Cys	Tyr	Ser	Gly	Gly	Ala	Val	Phe	Ser	Asp	Gly	Thr	Leu	Lys	Leu
	50					55				60					
Phe	Tyr	Thr	Gly	Asn	Arg	Lys	Ile	Asp	Gly	Lys	Arg	Arg	Ala	Thr	Gln
65				70				75					80		
Asn	Leu	Val	Glu	Val	Glu	Asp	Pro	Thr	Gly	Leu	Met	Gly	Gly	Ile	His
			85					90					95		
Arg	Arg	Ser	Pro	Lys	Asn	Pro	Leu	Ile	Asp	Gly	Pro	Ala	Ser	Gly	Phe
			100					105					110		
Thr	Pro	His	Tyr	Arg	Asp	Pro	Met	Ile	Ser	Pro	Asp	Gly	Asp	Gly	Trp
		115					120					125			
Lys	Met	Val	Leu	Gly	Ala	Gln	Arg	Glu	Asn	Leu	Thr	Gly	Ala	Ala	Val
	130					135					140				
Leu	Tyr	Arg	Ser	Ala	Asp	Leu	Glu	Asn	Trp	Glu	Phe	Ser	Gly	Glu	Ile
145				150				155					160		
Thr	Phe	Asp	Leu	Ser	Asp										
				165											

<210> 13  
 <211> 479  
 <212> DNA  
 <213> Corynebacterium thermoaminogenes

<220>

<221> CDS

<222> (1)..(477)

<400> 13

tac	tac	cag	cac	gat	cca	ggt	ttc	ccc	ttc	gca	cca	aag	cgc	acc	ggc
1				5				10					15		48
Tyr	Tyr	Gln	His	Asp	Pro	Gly	Phe	Pro	Phe	Ala	Pro	Lys	Arg	Thr	Gly

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1gg gct cac acc acc acg ccg ttg acc gga ccg cag cga ttg cag tgg	96
Trp Ala His Thr Thr Thr Pro Leu Thr Gly Pro Gln Arg Leu Gln Trp	
20 25 30	
acg cac ctg ccc gac gct ctt tac ccg gat gca tcc tat gac ctg gat	144
Thr His Leu Pro Asp Ala Leu Tyr Pro Asp Ala Ser Tyr Asp Leu Asp	
35 40 45	
gga tgc tat tcc ggt gga gcc gla ttt act gac ggc aca ctt aaa ctt	192
Gly Cys Tyr Ser Gly Gly Ala Val Phe Thr Asp Gly Thr Leu Lys Leu	
50 55 60	
ttc tac acc ggc aac cta aaa att gac ggc aag cgc cgc gcc acc caa	240
Phe Tyr Thr Gly Asn Leu Lys Ile Asp Gly Lys Arg Arg Ala Thr Gln	
65 70 75 80	
aac ctc gtc gaa gtc gag gac cca act ggg ctg atg ggc ggc att cat	288
Asn Leu Val Glu Val Glu Asp Pro Thr Gly Leu Met Gly Gly Ile His	
85 90 95	
cgc cgt tgc cct aaa aat ccg ctt atc gac gga ccc gcc agc ggt ttc	336
Arg Arg Ser Pro Lys Asn Pro Leu Ile Asp Gly Pro Ala Ser Gly Phe	
100 105 110	
aca ccc cat tac cgc gat ccc atg atc agc cct gat ggt gat ggt tgg	384
Thr Pro His Tyr Arg Asp Pro Met Ile Ser Pro Asp Gly Asp Gly Trp	
115 120 125	
aaa atg gtt ctt ggg gcc caa cgc gaa aac ctc acc ggt gca gcg gtt	432
Lys Met Val Leu Gly Ala Gln Arg Glu Asn Leu Thr Gly Ala Ala Val	
130 135 140	
cta tac cgc tgc aca gat ctt gaa aac tgg gaa ttc tcc ggt gaa at	479
Leu Tyr Arg Ser Thr Asp Leu Glu Asn Trp Glu Phe Ser Gly Glu	
145 150 155	

&lt;210&gt; 14

&lt;211&gt; 159

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 14

Tyr Tyr Gln His Asp Pro Gly Phe Pro Phe Ala Pro Lys Arg Thr Gly	
1 5 10 15	
Trp Ala His Thr Thr Thr Pro Leu Thr Gly Pro Gln Arg Leu Gln Trp	
20 25 30	
Thr His Leu Pro Asp Ala Leu Tyr Pro Asp Ala Ser Tyr Asp Leu Asp	
35 40 45	
Gly Cys Tyr Ser Gly Gly Ala Val Phe Thr Asp Gly Thr Leu Lys Leu	
50 55 60	
Phe Tyr Thr Gly Asn Leu Lys Ile Asp Gly Lys Arg Arg Ala Thr Gln	

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65		70		75		80									
Asn	Leu	Val	Glu	Val	Glu	Asp	Pro	Thr	Gly	Leu	Met	Gly	Gly	Ile	His
			85						90					95	
Arg	Arg	Ser	Pro	Lys	Asn	Pro	Leu	Ile	Asp	Gly	Pro	Ala	Ser	Gly	Phe
			100					105					110		
Thr	Pro	His	Tyr	Arg	Asp	Pro	Met	Ile	Ser	Pro	Asp	Gly	Asp	Gly	Trp
		115					120					125			
Lys	Met	Val	Leu	Gly	Ala	Gln	Arg	Glu	Asn	Leu	Thr	Gly	Ala	Ala	Val
	130					135				140					
Leu	Tyr	Arg	Ser	Thr	Asp	Leu	Glu	Asn	Trp	Glu	Phe	Ser	Gly	Glu	
145					150					155					

&lt;210&gt; 15

&lt;211&gt; 490

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 15

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atlttaalgg atattaicta tatlttatca atattatcct tatgcacctg aatggggacc 60
aatgcattgg ggacacgcac glagtaaaga lllagltlcal tgggaaacat taccgattgc 120
tllagaacct ggagaatgaag aagaaaaatg gtlgtttctc tggtaacaggt atagtcaaag 180
atgataagtt gtatltatll lalacagggt accattatta taatgacgat gatcccgatc 240
atlttllggca aaalcaaaal atggcittata glgaagaatgg cattcatttt caaaaatata 300
aacaaaatgc aatcatttct accccaccctg aagataaatc acatcacttc agagatccaa 360
aggtaatggga acatccaatgg cttattatta catgatagta ggtagltcaaa atgatagaga 420
attaggacgt attatcttat atcgttctga ggatttatag aggggaattc tggtcctgag 480
atcaatccaa                                     490

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&lt;210&gt; 16

&lt;211&gt; 4254

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (637)..(1362)

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1434)..(2315)

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (2432)..(3115)

27/123

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (3235).. (4065)

&lt;400&gt; 16

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tcacggcgcg cagattaccc agtggccgt agagacgcig atcggcattc tcacgcaccg 60
cgcaggtggt gaagacgatg agatcagggg tgcacccctc ccccgccgcg gtgtaacgg 120
ccctctcgag cagaccggag agacgctcgg aatcgtggac gtltatcigg cagccgaagg 180
tacgcaccctc ataggctcgg gcagtggtgc cctcccggtt ccccgcgcc gggagggtgt 240
cggcggggtg gtccgggtgg gatggaagg tgttcatcig gtgggtatca atctgtcgcg 300
tcacgggagg taattgtatc ggccgcgggc acccigacat aaacgtccga tccagaggaa 360
cgcaaccccg tggagtgctg cagccatgca ggttgggcaa caccgtaacg gaacctagca 420
gagtggtagg attgacttca cttctttac ctattgagct attgataaaa tccgggcgga 480
aatggaaatc acccccacaa atcaccccaa ctgacctgtg gaaagggcga gaaatccagg 540
gaaattcatt tcaaaatgga ctcaatcaca ggaattaccc cacatgacct aacattctt 600
tatgtatcc ccatgacgca gaccacaaat caccgc atg atc aag atg acg ggg 654
                                Met Ile Lys Met Thr Gly
                                1             5

gtg cag aag ttc ttc gat gac ttc cag gcc ctg acc gat atc aat ctt 702
Val Gln Lys Phe Phe Asp Asp Phe Gln Ala Leu Thr Asp Ile Asn Leu
                                10             15             20

gag gtc ccc gcg gga cag gtc gtt gtt gtt ctg gcc ccg tcc ggt tcc 750
Glu Val Pro Ala Gly Gln Val Val Val Val Leu Gly Pro Ser Gly Ser
                                25             30             35

gga aag tcg acg ctg tgc cgc acc atc aac cgc ctg gaa acc atc gag 798
Gly Lys Ser Thr Leu Cys Arg Thr Ile Asn Arg Leu Glu Thr Ile Glu
                                40             45             50

gag gga acc atc gag atc gat gga aaa ctg ctt ccg gag gag ggc aag 846
Glu Gly Thr Ile Glu Ile Asp Gly Lys Leu Leu Pro Glu Glu Gly Lys
                                55             60             65             70

gac ctg gcc aag atc cgt gcc gac gtg ggc atg gtg ttc cag tct ttc 894
Asp Leu Ala Lys Ile Arg Ala Asp Val Gly Met Val Phe Gln Ser Phe
                                75             80             85

aac ctg ttc ccc cac ctg acc atc aag gac aat gtc acc ctg gcc ccg 942
Asn Leu Phe Pro His Leu Thr Ile Lys Asp Asn Val Thr Leu Gly Pro
                                90             95             100

atg aag gtc cgg aag atg aag aag tcc gag gcc aat gag gtg gcc atg 990
Met Lys Val Arg Lys Met Lys Lys Ser Glu Ala Asn Glu Val Ala Met
                                105             110             115

aag ctg ttg gaa cgc gtc ggc atc gcc aac cag gcc gag aaa tac ccg 1038
Lys Leu Leu Glu Arg Val Gly Ile Ala Asn Gln Ala Glu Lys Tyr Pro
                                120             125             130

gca cag ctg tcg ggc ggg cag cag cag cgc gtg gcc atc gcc cgc gca 1086

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Ala Gln Leu Ser Gly Gly Gln Gln Gln Arg Val Ala Ile Ala Arg Ala	
135	140 145 150
ctg gcg atg aac ccc aag atc atg ctt ttc gac gaa cca acc tcc gcc	1134
Leu Ala Met Asn Pro Lys Ile Met Leu Phe Asp Glu Pro Thr Ser Ala	
155 160 165	
ctc gac ccc gag atg gtc aac gag gtt ctg gac gtc atg gcg agt ctg	1182
Leu Asp Pro Glu Met Val Asn Glu Val Leu Asp Val Met Ala Ser Leu	
170 175 180	
gcc aag gaa ggc atg acc atg gtg tgt gtc acc cac gag atg ggt ttc	1230
Ala Lys Glu Gly Met Thr Met Val Cys Val Thr His Glu Met Gly Phe	
185 190 195	
gca cgc agg gcc gca gac cgt gtg ctg ttc atg tct gac ggc gcc atc	1278
Ala Arg Arg Ala Ala Asp Arg Val Leu Phe Met Ser Asp Gly Ala Ile	
200 205 210	
gtc gag gac tcc gac ccg gag acc ttc ttc acc aat cca caa acc gac	1326
Val Glu Asp Ser Asp Pro Glu Thr Phe Phe Thr Asn Pro Gln Thr Asp	
215 220 225 230	
cgg gcg aag gat ttc ctg ggc aag atc ctc gcc cac tgacctcccc	1372
Arg Ala Lys Asp Phe Leu Gly Lys Ile Leu Ala His	
235 240	
tcacctctgtg tccaactccc ccgcctggcca aaatcagcga ccatgaccaa caggagcatc	1432
a atg tgc cac aaa cgc atg ttc acc cgt ctc gcc gca gcc acc agc gca	1481
Met Ser His Lys Arg Met Phe Thr Arg Leu Ala Ala Ala Thr Ser Ala	
245 250 255	
gct gtt ctc gcc ggc atc acc ctc acc gcc tgt ggt gat tcc gag ggt	1529
Ala Val Leu Ala Gly Ile Thr Leu Thr Ala Cys Gly Asp Ser Glu Gly	
260 265 270	
ggt gac ggt ctg ctc gcc gcc atc gaa aat ggc aat gtc acc atc ggc	1577
Gly Asp Gly Leu Leu Ala Ala Ile Glu Asn Gly Asn Val Thr Ile Gly	
275 280 285 290	
acc aag tac gat cag ccg ggt ctg gga ctg cgt aac ccg gac aat tcc	1625
Thr Lys Tyr Asp Gln Pro Gly Leu Gly Leu Arg Asn Pro Asp Asn Ser	
295 300 305	
atg agc gga ctg gat gtc gac gtc gcg cag tac gtg gtc aac tcc atc	1673
Met Ser Gly Leu Asp Val Asp Val Ala Gln Tyr Val Val Asn Ser Ile	
310 315 320	
gcc gat gac aac ggt tgg gat cac ccc acc gtg gaa tgg cgc gag acc	1721
Ala Asp Asp Asn Gly Trp Asp His Pro Thr Val Glu Trp Arg Glu Thr	
325 330 335	
ccc tcc gcc cag cgc gag acc ctc atc cag aac ggt gag gtg gat atg	1769
Pro Ser Ala Gln Arg Glu Thr Leu Ile Gln Asn Gly Glu Val Asp Met	
340 345 350	
atc gcc gca acc tac tcc atc aac ccc gga cgc tcc gaa tgc gtg aac	1817

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Ile	Ala	Ala	Thr	Tyr	Ser	Ile	Asn	Pro	Gly	Arg	Ser	Glu	Ser	Val	Asn		
355					360					365					370		
ttc	ggt	gga	cca	tac	ctc	ctc	acc	cac	cag	gcc	ctc	ctg	gtc	cgc	gag	1865	
Phe	Gly	Gly	Pro	Tyr	Leu	Leu	Thr	His	Gln	Ala	Leu	Leu	Val	Arg	Glu		
				375					380					385			
gac	gat	gac	cgc	atc	cag	acc	ctc	gag	gac	ctc	gat	gac	ggc	ctg	atc	1913	
Asp	Asp	Asp	Arg	Ile	Gln	Thr	Leu	Glu	Asp	Leu	Asp	Asp	Gly	Leu	Ile		
			390					395					400				
ctg	tgt	tcc	gtt	acc	gga	tcc	acc	ccc	gcc	cag	aag	gtc	aag	gat	gtc	1961	
Leu	Cys	Ser	Val	Thr	Gly	Ser	Thr	Pro	Ala	Gln	Lys	Val	Lys	Asp	Val		
		405					410					415					
ctc	ccc	ggc	gtc	cag	ctg	cag	gaa	tac	gac	acc	tac	tcc	tcc	tgt	gtg	2009	
Leu	Pro	Gly	Val	Gln	Leu	Gln	Glu	Tyr	Asp	Thr	Tyr	Ser	Ser	Cys	Val		
	420					425					430						
gag	gca	ctg	agc	cag	ggc	aac	gtc	gat	gca	atg	acc	acc	gac	gcc	acc	2057	
Glu	Ala	Leu	Ser	Gln	Gly	Asn	Val	Asp	Ala	Met	Thr	Thr	Asp	Ala	Thr		
435					440				445					450			
atc	ctc	ttc	ggc	tac	gcg	cag	cag	cgc	gaa	ggt	gaa	ttc	cgc	gtc	gtg	2105	
Ile	Leu	Phe	Gly	Tyr	Ala	Gln	Gln	Arg	Glu	Gly	Glu	Phe	Arg	Val	Val		
			455					460					465				
gag	atg	gaa	cag	gac	ggc	gag	ccg	ttc	acc	aat	gag	tac	tac	ggc	atc	2153	
Glu	Met	Glu	Gln	Asp	Gly	Glu	Pro	Phe	Thr	Asn	Glu	Tyr	Tyr	Gly	Ile		
		470					475				480						
ggt	atc	acc	aag	gat	gac	acc	gaa	gcc	acc	gat	gcg	atc	aac	gca	gcg	2201	
Gly	Ile	Thr	Lys	Asp	Asp	Thr	Glu	Ala	Thr	Asp	Ala	Ile	Asn	Ala	Ala		
	485					490			495								
tig	gag	cgt	atg	tac	gcc	gac	ggt	tcc	ttc	cag	cgt	ttc	ctc	acc	gag	2249	
Leu	Glu	Arg	Met	Tyr	Ala	Asp	Gly	Ser	Phe	Gln	Arg	Phe	Leu	Thr	Glu		
	500					505			510								
aac	ctc	ggc	gag	gat	tcc	cag	gtt	gtc	cag	gag	ggc	acc	ccg	ggt	gac	2297	
Asn	Leu	Gly	Glu	Asp	Ser	Gln	Val	Val	Gln	Glu	Gly	Thr	Pro	Gly	Asp		
515					520				525				530				
ctc	tcc	ttc	ctg	gac	gag	tgacc	tgacg	ggg	ccgaacg	ccc	gatgagc					2345	
Leu	Ser	Phe	Leu	Asp	Glu												
			535														
atgcgtggcc	cccgcattccc	ggggtgccac	gcatcatcac	tttcaccact	gatccccctac											2405	
cglttccctac	cgaggagaaaa	ttccccc	atg agt	aca ita	tgg gcg	gat ctg	ggt									2458	
			Met	Ser	Thr	Leu	Trp	Ala	Asp	Leu	Gly						
						540					545						
ccg	tca	ctc	cta	ccc	gca	ttc	tgg	gtg	aca	atc	caa	ctc	acc	gtc	tat	2506	
Pro	Ser	Leu	Leu	Pro	Ala	Phe	Trp	Val	Thr	Ile	Gln	Leu	Thr	Val	Tyr		
				550					555				560				
tcc	gcc	atc	gga	tcc	atg	atc	ctc	ggt	acc	atc	ctc	acc	gcc	atg	agg	2554	



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Ser	Ala	Ile	Gly	Ser	Met	Ile	Leu	Gly	Thr	Ile	Leu	Thr	Ala	Met	Arg	
			565			570						575				
gtg	icc	ccg	gtg	aag	atc	ctg	cgc	agc	ata	icc	acc	gcc	tac	atc	aac	2602
Val	Ser	Pro	Val	Lys	Ile	Leu	Arg	Ser	Ile	Ser	Thr	Ala	Tyr	Ile	Asn	
			580			585						590				
acg	gtc	cgt	aac	acc	cca	ctg	acc	ctg	gtg	atc	ctg	ttc	tgt	tcg	ttc	2650
Thr	Val	Arg	Asn	Thr	Pro	Leu	Thr	Leu	Val	Ile	Leu	Phe	Cys	Ser	Phe	
			595			600						605				
ggc	ctg	tat	cag	aat	ctc	ggt	ctc	acc	ctc	gcc	ggt	cgc	gac	agt	tcg	2698
Gly	Leu	Tyr	Gln	Asn	Leu	Gly	Leu	Thr	Leu	Ala	Gly	Arg	Asp	Ser	Ser	
610			615						620			625				
acc	ttt	ctg	gcc	gat	aac	aac	ttc	cgg	ctc	gcg	gtg	ctc	gga	ttc	atc	2746
Thr	Phe	Leu	Ala	Asp	Asn	Asn	Phe	Arg	Leu	Ala	Val	Leu	Gly	Phe	Ile	
			630						635			640				
ctg	tac	acc	tcg	gcc	ttc	gtt	gcg	gaa	tca	ctc	cgg	tca	ggc	atc	aac	2794
Leu	Tyr	Thr	Ser	Ala	Phe	Val	Ala	Glu	Ser	Leu	Arg	Ser	Gly	Ile	Asn	
			645						650			655				
acc	gtg	cac	ttc	ggg	cag	gcg	gag	gcc	gcc	cgg	tcg	ctg	gga	ctc	ggt	2842
Thr	Val	His	Phe	Gly	Gln	Ala	Glu	Ala	Ala	Arg	Ser	Leu	Gly	Leu	Gly	
			660			665						670				
ttc	agt	gac	atc	ttc	cgg	tcg	atc	atc	ttc	ccc	cag	gcg	gtg	cgt	gcc	2890
Phe	Ser	Asp	Ile	Phe	Arg	Ser	Ile	Ile	Phe	Pro	Gln	Ala	Val	Arg	Ala	
675			680						685							
gcc	atc	atc	ccg	ctg	ggc	aac	acc	ctc	atc	gcc	ctg	acc	aag	aac	acc	2938
Ala	Ile	Ile	Pro	Leu	Gly	Asn	Thr	Leu	Ile	Ala	Leu	Thr	Lys	Asn	Thr	
690			695						700			705				
acg	atc	gcg	tcg	gtg	atc	ggc	gtc	ggt	gag	gcc	tcg	ctg	ctg	atg	aag	2986
Thr	Ile	Ala	Ser	Val	Ile	Gly	Val	Gly	Glu	Ala	Ser	Leu	Leu	Met	Lys	
			710						715			720				
tcg	acg	att	gaa	aat	cat	gcc	aac	atg	ctc	ttc	gtc	gtg	ttc	gcc	atc	3034
Ser	Thr	Ile	Glu	Asn	His	Ala	Asn	Met	Leu	Phe	Val	Val	Phe	Ala	Ile	
			725						730			735				
ttc	gcc	gtc	ggc	ttc	atg	atc	ctc	acc	ctc	ccc	atg	ggc	ctg	ggg	ctt	3082
Phe	Ala	Val	Gly	Phe	Met	Ile	Leu	Thr	Leu	Pro	Met	Gly	Leu	Gly	Leu	
740						745						750				
gga	aaa	ctc	gct	gag	aaa	atg	gcg	gtg	aag	aaa	taatgttcctc	ctccgtacgc				3135
Gly	Lys	Leu	Ala	Glu	Lys	Met	Ala	Val	Lys	Lys						
755						760										
gcaacagttcc tctacgacgc ccccggtccc cggggacgca ggtccaacac catcatcacc																3195
atcgccacca ccttggtggc agtggccgtc ctgttctgg gtg ggc agt gtt ctc																3249
Val Gly Ser Val Leu																
765																
cag gaa aac ggc cag ttg gac ggc gac aaa tgg acc ccg ttc ctc gat																3297

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Gln Glu Asn Gly Gln Leu Asp Gly Asp Lys Trp Thr Pro Phe Leu Asp	
770 775 780 785	
ccc cag acc tgg acc acc tat ctt ctg ccc ggc ctg tgg gga acc ctg	3345
Pro Gln Thr Trp Thr Thr Tyr Leu Leu Pro Gly Leu Trp Gly Thr Leu	
790 795 800	
aag gca gcg gtg gcc tcc atc ctt ctc gcg ctg atc atg ggc acc ctg	3393
Lys Ala Ala Val Ala Ser Ile Leu Leu Ala Leu Ile Met Gly Thr Leu	
805 810 815	
ctc ggg ctc gga cgc atc tcc gaa atc cgg ctc ctg cgc tgg ttc tgc	3441
Leu Gly Leu Gly Arg Ile Ser Glu Ile Arg Leu Leu Arg Trp Phe Cys	
820 825 830	
ggg atc atc atc gag acc ttc cgt gcc atc ccg gtg ctg atc ctc atg	3489
Gly Ile Ile Ile Glu Thr Phe Arg Ala Ile Pro Val Leu Ile Leu Met	
835 840 845	
atc ttc gcc tat cag ttg ttc gcc cgt tac cag ctc gtt cca tca cgc	3537
Ile Phe Ala Tyr Gln Leu Phe Ala Arg Tyr Gln Leu Val Pro Ser Arg	
850 855 860 865	
cag ctg gcc ttc gcc gcg gtg gtc ttc ggt ctc acc atg tac aac ggc	3585
Gln Leu Ala Phe Ala Ala Val Val Phe Gly Leu Thr Met Tyr Asn Gly	
870 875 880	
tcc gtc atc gcc gag atc ctt aga tcc ggt atc gcc tcc ctg ccg aag	3633
Ser Val Ile Ala Glu Ile Leu Arg Ser Gly Ile Ala Ser Leu Pro Lys	
885 890 895	
gga cag cgt gag gcg gcg atc gcc ctg ggc atg tca acc cgc cag acc	3681
Gly Gln Arg Glu Ala Ala Ile Ala Leu Gly Met Ser Thr Arg Gln Thr	
900 905 910	
acc tgg tcc atc ctg ctc ccc cag gcg gtg gca gcg atg ctg ccc gcc	3729
Thr Trp Ser Ile Leu Leu Pro Gln Ala Val Ala Ala Met Leu Pro Ala	
915 920 925	
ctg atc gcg cag atg gtc atc gcg ctg aag gac tcc gcc ctc ggt tac	3777
Leu Ile Ala Gln Met Val Ile Ala Leu Lys Asp Ser Ala Leu Gly Tyr	
930 935 940 945	
cag atc ggt tat atc gag gtg gla cgc tcc ggt atc cag tcc gca tcc	3825
Gln Ile Gly Tyr Ile Glu Val Val Arg Ser Gly Ile Gln Ser Ala Ser	
950 955 960	
gtc aac cgg aac tac ctg gct gcc ctc gcg gtg gtc gcg gtc atc atg	3873
Val Asn Arg Asn Tyr Leu Ala Ala Leu Ala Val Val Ala Val Ile Met	
965 970 975	
atc ctg atc aac ttc gca ctg acc gca ctg gca gag cgt atc cag cgt	3921
Ile Leu Ile Asn Phe Ala Leu Thr Ala Leu Ala Glu Arg Ile Gln Arg	
980 985 990	
cag ctg cgt gcc gga cgt gcc cgc agg aac att gtg gca aag gtg ccc	3969
Gln Leu Arg Ala Gly Arg Ala Arg Arg Asn Ile Val Ala Lys Val Pro	

<400> 17																
Met	Ile	Lys	Met	Thr	Gly	Val	Gln	Lys	Phe	Phe	Asp	Asp	Phe	Gln	Ala	
1				5					10					15		
Leu	Thr	Asp	Ile	Asn	Leu	Glu	Val	Pro	Ala	Gly	Gln	Val	Val	Val	Val	
			20					25					30			
Leu	Gly	Pro	Ser	Gly	Ser	Gly	Lys	Ser	Thr	Leu	Cys	Arg	Thr	Ile	Asn	
		35					40					45				
Arg	Leu	Glu	Thr	Ile	Glu	Glu	Gly	Thr	Ile	Glu	Ile	Asp	Gly	Lys	Leu	
	50					55					60					
Leu	Pro	Glu	Glu	Gly	Lys	Asp	Leu	Ala	Lys	Ile	Arg	Ala	Asp	Val	Gly	
65					70					75					80	
Met	Val	Phe	Gln	Ser	Phe	Asn	Leu	Phe	Pro	His	Leu	Thr	Ile	Lys	Asp	
			85					90						95		
Asn	Val	Thr	Leu	Gly	Pro	Met	Lys	Val	Arg	Lys	Met	Lys	Lys	Ser	Glu	
			100					105					110			
Ala	Asn	Glu	Val	Ala	Met	Lys	Leu	Leu	Glu	Arg	Val	Gly	Ile	Ala	Asn	
		115					120					125				
Gln	Ala	Glu	Lys	Tyr	Pro	Ala	Gln	Leu	Ser	Gly	Gly	Gln	Gln	Gln	Arg	
	130					135				140						
Val	Ala	Ile	Ala	Arg	Ala	Leu	Ala	Met	Asn	Pro	Lys	Ile	Met	Leu	Phe	
145				150					155					160		
Asp	Glu	Pro	Thr	Ser	Ala	Leu	Asp	Pro	Glu	Met	Val	Asn	Glu	Val	Leu	
			165					170					175			
Asp	Val	Met	Ala	Ser	Leu	Ala	Lys	Glu	Gly	Met	Thr	Met	Val	Cys	Val	
		180					185					190				
Thr	His	Glu	Met	Gly	Phe	Ala	Arg	Arg	Ala	Ala	Asp	Arg	Val	Leu	Phe	

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	195		200		205	
Met	Ser Asp Gly Ala Ile Val Glu Asp Ser Asp Pro Glu Thr Phe Phe					
	210		215		220	
Thr Asn Pro Gln Thr Asp Arg Ala Lys Asp Phe Leu Gly Lys Ile Leu						
225		230		235		240
Ala His						

&lt;210&gt; 18

&lt;211&gt; 294

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 18

Met	Ser His Lys Arg Met Phe Thr Arg Leu Ala Ala Ala Thr Ser Ala	
1	5	10 15
Ala Val Leu Ala Gly Ile Thr Leu Thr Ala Cys Gly Asp Ser Glu Gly		
20	25	30
Gly Asp Gly Leu Leu Ala Ala Ile Glu Asn Gly Asn Val Thr Ile Gly		
35	40	45
Thr Lys Tyr Asp Gln Pro Gly Leu Gly Leu Arg Asn Pro Asp Asn Ser		
50	55	60
Met Ser Gly Leu Asp Val Asp Val Ala Gln Tyr Val Val Asn Ser Ile		
65	70	75 80
Ala Asp Asp Asn Gly Trp Asp His Pro Thr Val Glu Trp Arg Glu Thr		
85	90	95
Pro Ser Ala Gln Arg Glu Thr Leu Ile Gln Asn Gly Glu Val Asp Met		
100	105	110
Ile Ala Ala Thr Tyr Ser Ile Asn Pro Gly Arg Ser Glu Ser Val Asn		
115	120	125
Phe Gly Gly Pro Tyr Leu Leu Thr His Gln Ala Leu Leu Val Arg Glu		
130	135	140
Asp Asp Asp Arg Ile Gln Thr Leu Glu Asp Leu Asp Asp Gly Leu Ile		
145	150	155 160
Leu Cys Ser Val Thr Gly Ser Thr Pro Ala Gln Lys Val Lys Asp Val		
165	170	175
Leu Pro Gly Val Gln Leu Gln Glu Tyr Asp Thr Tyr Ser Ser Cys Val		
180	185	190
Glu Ala Leu Ser Gln Gly Asn Val Asp Ala Met Thr Thr Asp Ala Thr		
195	200	205
Ile Leu Phe Gly Tyr Ala Gln Arg Glu Gly Glu Phe Arg Val Val		
210	215	220
Glu Met Glu Gln Asp Gly Glu Pro Phe Thr Asn Glu Tyr Tyr Gly Ile		
225	230	235 240

Met	Ser	Thr	Leu	Trp	Ala	Asp	Leu	Gly	Pro	Ser	Leu	Leu	Pro	Ala	Phe
1				5				10					15		
Trp	Val	Thr	Ile	Gln	Leu	Thr	Val	Tyr	Ser	Ala	Ile	Gly	Ser	Met	Ile
			20					25					30		
Leu	Gly	Thr	Ile	Leu	Thr	Ala	Met	Arg	Val	Ser	Pro	Val	Lys	Ile	Leu
		35					40					45			
Arg	Ser	Ile	Ser	Thr	Ala	Tyr	Ile	Asn	Thr	Val	Arg	Asn	Thr	Pro	Leu
	50					55					60				
Thr	Leu	Val	Ile	Leu	Phe	Cys	Ser	Phe	Gly	Leu	Tyr	Gln	Asn	Leu	Gly
65					70					75					80
Leu	Thr	Leu	Ala	Gly	Arg	Asp	Ser	Ser	Thr	Phe	Leu	Ala	Asp	Asn	Asn
				85					90					95	
Phe	Arg	Leu	Ala	Val	Leu	Gly	Phe	Ile	Leu	Tyr	Thr	Ser	Ala	Phe	Val
			100					105					110		
Ala	Glu	Ser	Leu	Arg	Ser	Gly	Ile	Asn	Thr	Val	His	Phe	Gly	Gln	Ala
		115					120					125			
Glu	Ala	Ala	Arg	Ser	Leu	Gly	Leu	Gly	Phe	Ser	Asp	Ile	Phe	Arg	Ser
	130					135					140				
Ile	Ile	Phe	Pro	Gln	Ala	Val	Arg	Ala	Ala	Ile	Ile	Pro	Leu	Gly	Asn
145					150					155					160
Thr	Leu	Ile	Ala	Leu	Thr	Lys	Asn	Thr	Thr	Ile	Ala	Ser	Val	Ile	Gly
				165					170					175	
Val	Gly	Glu	Ala	Ser	Leu	Leu	Met	Lys	Ser	Thr	Ile	Glu	Asn	His	Ala
			180					185				190			
Asn	Met	Leu	Phe	Val	Val	Phe	Ala	Ile	Phe	Ala	Val	Gly	Phe	Met	Ile
		195					200					205			
Leu	Thr	Leu	Pro	Met	Gly	Leu	Gly	Leu	Gly	Lys	Leu	Ala	Glu	Lys	Met
	210					215						220			

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Ala Val Lys Lys

225

&lt;210&gt; 20

&lt;211&gt; 277

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 20

Val	Gly	Ser	Val	Leu	Gln	Glu	Asn	Gly	Gln	Leu	Asp	Gly	Asp	Lys	Trp
1				5					10					15	
Thr	Pro	Phe	Leu	Asp	Pro	Gln	Thr	Trp	Thr	Thr	Tyr	Leu	Leu	Pro	Gly
			20					25					30		
Leu	Trp	Gly	Thr	Leu	Lys	Ala	Ala	Val	Ala	Ser	Ile	Leu	Leu	Ala	Leu
		35					40					45			
Ile	Met	Gly	Thr	Leu	Leu	Gly	Leu	Gly	Arg	Ile	Ser	Glu	Ile	Arg	Leu
	50					55				60					
Leu	Arg	Trp	Phe	Cys	Gly	Ile	Ile	Ile	Glu	Thr	Phe	Arg	Ala	Ile	Pro
65					70				75					80	
Val	Leu	Ile	Leu	Met	Ile	Phe	Ala	Tyr	Gln	Leu	Phe	Ala	Arg	Tyr	Gln
				85					90					95	
Leu	Val	Pro	Ser	Arg	Gln	Leu	Ala	Phe	Ala	Ala	Val	Val	Phe	Gly	Leu
		100					105					110			
Thr	Met	Tyr	Asn	Gly	Ser	Val	Ile	Ala	Glu	Ile	Leu	Arg	Ser	Gly	Ile
	115						120					125			
Ala	Ser	Leu	Pro	Lys	Gly	Gln	Arg	Glu	Ala	Ala	Ile	Ala	Leu	Gly	Met
130						135					140				
Ser	Thr	Arg	Gln	Thr	Thr	Trp	Ser	Ile	Leu	Leu	Pro	Gln	Ala	Val	Ala
145					150				155					160	
Ala	Met	Leu	Pro	Ala	Leu	Ile	Ala	Gln	Met	Val	Ile	Ala	Leu	Lys	Asp
				165					170					175	
Ser	Ala	Leu	Gly	Tyr	Gln	Ile	Gly	Tyr	Ile	Glu	Val	Val	Arg	Ser	Gly
		180						185					190		
Ile	Gln	Ser	Ala	Ser	Val	Asn	Arg	Asn	Tyr	Leu	Ala	Ala	Leu	Ala	Val
	195						200						205		
Val	Ala	Val	Ile	Met	Ile	Leu	Ile	Asn	Phe	Ala	Leu	Thr	Ala	Leu	Ala
	210					215					220				
Glu	Arg	Ile	Gln	Arg	Gln	Leu	Arg	Ala	Gly	Arg	Ala	Arg	Arg	Asn	Ile
225					230				235					240	
Val	Ala	Lys	Val	Pro	Glu	Glu	Pro	Asp	Gln	Gly	Leu	Asp	Thr	Lys	Asp
				245					250					255	
Asn	Val	Asn	Val	Asp	Trp	His	Asp	Pro	Asp	Tyr	Lys	Glu	Val	Lys	His
			260					265						270	

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Pro Gly Pro Ser Phe

275

&lt;210&gt; 21

&lt;211&gt; 3598

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (454)..(3222)

&lt;400&gt; 21

agcacggcca aacatgagag aaacttcaca ttttgaattt cccctttcct gcataatggaa 60  
 aaccgcccgt gacacccctg ccatttgggc agctcccccc acctcaccat gtccacattt 120  
 tccataaigt ggccigttaac accttgggc tcaaggcttc caegccccac cgggaccctc 180  
 atcagcagggt gaaacagacc ctcttgaat gctttgttaa aaagaaccgc cctttgtgcg 240  
 taiccttgtg tcaattgtgc ggcacatgcc accagcttcc ctacagattg aacacggtcg 300  
 ggaaatcctc cccggatacc ctgcacgccc caccctccac accgacaccg gcggggaggg 360  
 ccgggcacgt tttcagctgc gggatgatga agcggctgcc ggtcccccggt tgcataaac 420  
 gaaatgaaaa acattccaac aggagggtgtg gaa atg gcc gat caa gca aaa ctt 474

Met Ala Asp Gln Ala Lys Leu

1

5

ggt ggc aaa ccc aca gat gac acc aac ttc gcg atg atc cgt gat ggc 522  
 Gly Gly Lys Pro Thr Asp Asp Thr Asn Phe Ala Met Ile Arg Asp Gly

10

15

20

gtt gca tct tat ttg aac gac tcc gac ccg gag gag acc aag gag tgg 570  
 Val Ala Ser Tyr Leu Asn Asp Ser Asp Pro Glu Glu Thr Lys Glu Trp

25

30

35

atg gac tcc cta gac ggt cta ctg cag gat tcc tct ccg gag cgc gcc 618  
 Met Asp Ser Leu Asp Gly Leu Leu Gln Asp Ser Ser Pro Glu Arg Ala

40

45

50

55

cgt tac ctg atg ctg cgc ctg ctg gag cgg gca tcc gcc aag cgt gtc 666  
 Arg Tyr Leu Met Leu Arg Leu Leu Glu Arg Ala Ser Ala Lys Arg Val

60

65

70

cca ctg ccc ccg atg acg tcc acc gat tac gtc aac acc atc ccc aca 714  
 Pro Leu Pro Pro Met Thr Ser Thr Asp Tyr Val Asn Thr Ile Pro Thr

75

80

85

tcc atg gag ccc gat ttc ccg ggt gat gag gag atg gag aag cgc tac 762  
 Ser Met Glu Pro Asp Phe Pro Gly Asp Glu Glu Met Glu Lys Arg Tyr

90

95

100

cgc cgc tgg atg cgc tgg aac gcc gcc atc atg gtg cac cgt gcc cag 810  
 Arg Arg Trp Met Arg Trp Asn Ala Ala Ile Met Val His Arg Ala Gln

105

110

115

cgc ccg gga atc ggt gtg ggt ggg cac atc tcc acc tac gcc ggc gcc 858

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Arg	Pro	Gly	Ile	Gly	Val	Gly	Gly	His	Ile	Ser	Thr	Tyr	Ala	Gly	Ala		
120					125					130					135		
gcc	cca	ctc	tac	gag	gtc	ggt	ttc	aac	cac	ttc	ttc	cgc	ggc	aag	gac	906	
Ala	Pro	Leu	Tyr	Glu	Val	Gly	Phe	Asn	His	Phe	Phe	Arg	Gly	Lys	Asp		
				140						145					150		
cac	ccg	ggt	ggc	ggt	gac	cag	gtc	ttc	ttc	cag	ggt	cac	gcc	tcc	ccg	954	
His	Pro	Gly	Gly	Gly	Asp	Gln	Val	Phe	Phe	Gln	Gly	His	Ala	Ser	Pro		
			155							160					165		
ggc	atg	tac	gcc	cgc	gcc	ttc	ctc	gag	ggc	cgt	ctc	acc	gag	agc	gat	1002	
Gly	Met	Tyr	Ala	Arg	Ala	Phe	Leu	Glu	Gly	Arg	Leu	Thr	Glu	Ser	Asp		
	170							175				180					
ctg	gac	agc	ttc	cgc	cag	gag	gtc	tcc	tac	gaa	ggt	ggt	ggc	atc	ccg	1050	
Leu	Asp	Ser	Phe	Arg	Gln	Glu	Val	Ser	Tyr	Glu	Gly	Gly	Gly	Gly	Ile	Pro	
	185						190					195					
tcc	tac	ccg	cac	ccg	cac	ggc	atg	ccg	gac	ttc	tgg	gag	ttc	ccg	acc	1098	
Ser	Tyr	Pro	His	Pro	His	Gly	Met	Pro	Asp	Phe	Trp	Glu	Phe	Pro	Thr		
200					205					210					215		
gtg	tcc	atg	ggc	ctc	ggg	ccc	atg	gat	gcc	atc	tac	cag	gcg	cgc	ttc	1146	
Val	Ser	Met	Gly	Leu	Gly	Pro	Met	Asp	Ala	Ile	Tyr	Gln	Ala	Arg	Phe		
				220						225					230		
aac	cgc	tac	ctg	cac	aac	cgt	ggc	atc	aag	gac	acc	tgc	gag	cag	cac	1194	
Asn	Arg	Tyr	Leu	His	Asn	Arg	Gly	Ile	Lys	Asp	Thr	Ser	Glu	Gln	His		
			235							240					245		
gtc	igg	gca	ttc	ctc	ggt	gac	ggc	gag	atg	gat	gag	ccg	gag	tcc	cgt	1242	
Val	Trp	Ala	Phe	Leu	Gly	Asp	Gly	Glu	Met	Asp	Glu	Pro	Glu	Ser	Arg		
	250						255					260					
ggt	ctc	atc	cac	cag	gct	gcg	ctg	aac	aac	ctg	gac	aac	ctc	acc	ttc	1290	
Gly	Leu	Ile	His	Gln	Ala	Ala	Leu	Asn	Asn	Leu	Asp	Asn	Leu	Thr	Phe		
	265						270					275					
gtg	atc	aac	tgc	aac	ctg	cag	cgt	ctt	gat	ggc	ccg	gtc	cgc	ggt	aac	1338	
Val	Ile	Asn	Cys	Asn	Leu	Gln	Arg	Leu	Asp	Gly	Pro	Val	Arg	Gly	Asn		
280					285					290					295		
acc	aag	atc	atc	cag	gaa	ctc	gag	tcc	ttc	ttc	cgt	ggt	gcc	ggc	tgg	1386	
Thr	Lys	Ile	Ile	Gln	Glu	Leu	Glu	Ser	Phe	Phe	Arg	Gly	Ala	Gly	Trp		
				300						305					310		
tcc	gtc	atc	aag	gtc	atc	igg	ggc	cgt	gag	igg	gat	gaa	ctg	ctg	gag	1434	
Ser	Val	Ile	Lys	Val	Ile	Trp	Gly	Arg	Glu	Trp	Asp	Glu	Leu	Leu	Glu		
			315							320					325		
aag	gac	cag	gac	ggt	gct	ctt	gtc	gag	gtc	atg	aac	aac	acc	tcc	gac	1482	
Lys	Asp	Gln	Asp	Gly	Ala	Leu	Val	Glu	Val	Met	Asn	Asn	Thr	Ser	Asp		
	330						335					340					
ggt	gac	tac	cag	acc	ttc	aag	gcc	aat	gac	ggt	gcc	tac	gtc	cgt	gag	1530	
Gly	Asp	Tyr	Gln	Thr	Phe	Lys	Ala	Asn	Asp	Gly	Ala	Tyr	Val	Arg	Glu		



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345	350	355	
cac ttc ttc ggc cgt gac ccc cgc acc ctc aag ctc gtc gag gac atg			1578
His Phe Phe Gly Arg Asp Pro Arg Thr Leu Lys Leu Val Glu Asp Met			
360	365	370	375
acc gac gag gag atc tgg aag ctg ccc cgt ggt ggc cat gac tac cgt			1626
Thr Asp Glu Glu Ile Trp Lys Leu Pro Arg Gly Gly His Asp Tyr Arg			
380	385	390	
aag gtc tac gcc gcc tac aag cgt gcg ctg gag acc aag gac cgc ccg			1674
Lys Val Tyr Ala Ala Tyr Lys Arg Ala Leu Glu Thr Lys Asp Arg Pro			
395	400	405	
acc gtc att ctc gcc cat acc atc aag ggc tac ggc ctg ggc cac aac			1722
Thr Val Ile Leu Ala His Thr Ile Lys Gly Tyr Gly Leu Gly His Asn			
410	415	420	
ttc gag ggc cgc aac gcg acc cac cag atg aag aag ctg acc ctg gat			1770
Phe Glu Gly Arg Asn Ala Thr His Gln Met Lys Lys Leu Thr Leu Asp			
425	430	435	
gac ctg aag ctg ttc cgt gac aag cag ggt ctg ccc atc acc gat gag			1818
Asp Leu Lys Leu Phe Arg Asp Lys Gln Gly Leu Pro Ile Thr Asp Glu			
440	445	450	455
gag ctg gag aag gat ccc tac ctg cct ccg tac tac cac ccg ggt gag			1866
Glu Leu Glu Lys Asp Pro Tyr Leu Pro Pro Tyr Tyr His Pro Gly Glu			
460	465	470	
gac gca ccg gag atc aag tac atg aag gag cgt cgc cag gcg ctc ggt			1914
Asp Ala Pro Glu Ile Lys Tyr Met Lys Glu Arg Arg Gln Ala Leu Gly			
475	480	485	
ggt ttc ctg ccg gag cgc cgt gag aag tac gag cca ctg cag gtt ccc			1962
Gly Phe Leu Pro Glu Arg Arg Glu Lys Tyr Glu Pro Leu Gln Val Pro			
490	495	500	
ccg ctg gac aag ctg cgg tcc gtg cgc aag ggt tcc ggc aag cag cag			2010
Pro Leu Asp Lys Leu Arg Ser Val Arg Lys Gly Ser Gly Lys Gln Gln			
505	510	515	
gtg gcc acc acc atg gcc acg gtg cgt acc ttc aag gaa ctc atg cgg			2058
Val Ala Thr Thr Met Ala Thr Val Arg Thr Phe Lys Glu Leu Met Arg			
520	525	530	535
gac aag aac ctg gcc gac cgc ttg gtc ccg atc atc ccg gat gag gcc			2106
Asp Lys Asn Leu Ala Asp Arg Leu Val Pro Ile Ile Pro Asp Glu Ala			
540	545	550	
cgc acc ttc ggc ctg gac tcc tgg ttc ccg acc ctg aaa atc tac aac			2154
Arg Thr Phe Gly Leu Asp Ser Trp Phe Pro Thr Leu Lys Ile Tyr Asn			
555	560	565	
ccg cac ggt cag aac tac gtg ccg gtc gac cat gac ctc atg ctg tcc			2202
Pro His Gly Gln Asn Tyr Val Pro Val Asp His Asp Leu Met Leu Ser			
570	575	580	

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tac cgt gag gcc aag gac ggc cag atc ctg cat gag ggc atc aac gag	2250
Tyr Arg Glu Ala Lys Asp Gly Gln Ile Leu His Glu Gly Ile Asn Glu	
585 590 595	
gcc ggt tcc gtg gca tct ttt atc gcc gcc gga acc tcc tac gcc acc	2298
Ala Gly Ser Val Ala Ser Phe Ile Ala Ala Gly Thr Ser Tyr Ala Thr	
600 605 610 615	
cat ggc gag gcc atg atc ccg ctg tac atc ttc tac tct atg ttc ggc	2346
His Gly Glu Ala Met Ile Pro Leu Tyr Ile Phe Tyr Ser Met Phe Gly	
620 625 630	
ttc cag cgc acc ggt gac ggc atc tgg gcc gca gcc gac cag atg acg	2394
Phe Gln Arg Thr Gly Asp Gly Ile Trp Ala Ala Ala Asp Gln Met Thr	
635 640 645	
cgt ggt ttc ctg ctg ggc gcc acc gcc ggt cgc acc acc ctg acc ggt	2442
Arg Gly Phe Leu Leu Gly Ala Thr Ala Gly Arg Thr Thr Leu Thr Gly	
650 655 660	
gag ggc ctg cag cac atg gat ggc cac tcc ccg atc ctg gcc tcc acc	2490
Glu Gly Leu Gln His Met Asp Gly His Ser Pro Ile Leu Ala Ser Thr	
665 670 675	
aac ccc ggt gtg gag acc tat gac ccg gcg ttc tcc tac gag atc gcg	2538
Asn Pro Gly Val Glu Thr Tyr Asp Pro Ala Phe Ser Tyr Glu Ile Ala	
680 685 690 695	
cac ctg gtc cac cgc ggc atc gac cgc atg tac gga ccg ggc aag ggt	2586
His Leu Val His Arg Gly Ile Asp Arg Met Tyr Gly Pro Gly Lys Gly	
700 705 710	
gag aat gtc atc tac tac ctg acc atc tac aac gag cca acc ccg cag	2634
Glu Asn Val Ile Tyr Tyr Leu Thr Ile Tyr Asn Glu Pro Thr Pro Gln	
715 720 725	
ccg gct gag cct gag gat ctg gac gtc gag ggc ctg cac aag ggc atc	2682
Pro Ala Glu Pro Glu Asp Leu Asp Val Glu Gly Leu His Lys Gly Ile	
730 735 740	
tac ctg tac gac aag gcc gcc gag ggt gag ggc cat gag gcc tct atc	2730
Tyr Leu Tyr Asp Lys Ala Ala Glu Gly Glu Gly His Glu Ala Ser Ile	
745 750 755	
ctg gcc tcc ggc atc ggc atg cag tgg gca ctg cgc gcc cgt gac atc	2778
Leu Ala Ser Gly Ile Gly Met Gln Trp Ala Leu Arg Ala Arg Asp Ile	
760 765 770 775	
ctc gcc gag gat tac ggc atc cgt gcc aac atc ttc tcc gcc acc tct	2826
Leu Ala Glu Asp Tyr Gly Ile Arg Ala Asn Ile Phe Ser Ala Thr Ser	
780 785 790	
tgg gtg gag ctg gcc cgc gac ggt gcc cgc cgt aac ctg gag gcg ctg	2874
Trp Val Glu Leu Ala Arg Asp Gly Ala Arg Arg Asn Leu Glu Ala Leu	
795 800 805	
cgc aac ccg ggt gcg gat gtc ggt gag gca ttc gtg acc acc cag ctg	2922

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Arg	Asn	Pro	Gly	Ala	Asp	Val	Gly	Glu	Ala	Phe	Val	Thr	Thr	Gln	Leu		
	810						815				820						
aag	aag	ggt	tcc	ggc	ccc	tac	gtc	gcg	gtg	tcc	gac	ttc	gcg	acc	gac	2970	
Lys	Lys	Gly	Ser	Gly	Pro	Tyr	Val	Ala	Val	Ser	Asp	Phe	Ala	Thr	Asp		
	825					830					835						
ctg	ccg	aac	cag	atc	cgc	gag	tgg	gtt	ccc	ggt	gac	tac	atc	gtc	ctc	3018	
Leu	Pro	Asn	Gln	Ile	Arg	Glu	Trp	Val	Pro	Gly	Asp	Tyr	Ile	Val	Leu		
840					845					850					855		
ggt	gcc	gac	ggc	ttc	ggt	ttc	tcc	gat	acc	cgt	ccg	gca	gcc	cgt	cgt	3066	
Gly	Ala	Asp	Gly	Phe	Gly	Phe	Ser	Asp	Thr	Arg	Pro	Ala	Ala	Arg	Arg		
			860						865					870			
tac	ttc	aac	atc	gac	gcc	gag	tcc	atc	gtc	gtg	gcg	gtc	ctg	cgc	ggc	3114	
Tyr	Phe	Asn	Ile	Asp	Ala	Glu	Ser	Ile	Val	Val	Ala	Val	Leu	Arg	Gly		
		875						880					885				
ctg	gtc	cgc	gag	ggt	gtc	atc	gat	gcc	tcc	gtg	gcg	gcg	cac	gcg	gct	3162	
Leu	Val	Arg	Glu	Gly	Val	Ile	Asp	Ala	Ser	Val	Ala	Ala	His	Ala	Ala		
	890					895					900						
gag	aag	tac	aag	ctg	tcc	gac	ccg	acg	gca	cca	cag	gtc	gat	ccg	gac	3210	
Glu	Lys	Tyr	Lys	Leu	Ser	Asp	Pro	Thr	Ala	Pro	Gln	Val	Asp	Pro	Asp		
	905					910					915						
gca	ccg	atc	gag	tagacc	tgt	tgtc	gac	gaa	aaacac	cccc	gccccctc	ac				3262	
Ala	Pro	Ile	Glu														
920																	
atgatgaggg	gggcgggggt	gtgctcgltt	acggcgggia	caggggggla	tcagcccagc	3322											
atcgccitai	cggagagcgt	cgcgcccttg	atcttggcga	attccctgcag	cagatcccgc	3382											
acggtgagct	tcgtcttcac	ctctgcgcig	gcctcataga	cgatccgtcc	ctcgtgcate	3442											
atgatgaggc	ggliacccag	gcggatagcc	tgltccalgt	tgigggtgac	calgagggig	3502											
gtcagtttgc	cgicctcgac	gatcttctcg	gicagggig	tgaccagtic	ggctcgctgg	3562											
gggtccaggg	cggcgggtig	ttcgtcgaga	agcatg			3598											

&lt;210&gt; 22

&lt;211&gt; 923

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 22

Met	Ala	Asp	Gln	Ala	Lys	Leu	Gly	Gly	Lys	Pro	Thr	Asp	Asp	Thr	Asn		
1				5					10					15			
Phe	Ala	Met	Ile	Arg	Asp	Gly	Val	Ala	Ser	Tyr	Leu	Asn	Asp	Ser	Asp		
		20						25					30				
Pro	Glu	Glu	Thr	Lys	Glu	Trp	Met	Asp	Ser	Leu	Asp	Gly	Leu	Leu	Gln		
	35					40						45					
Asp	Ser	Ser	Pro	Glu	Arg	Ala	Arg	Tyr	Leu	Met	Leu	Arg	Leu	Leu	Glu		

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50		55		60
Arg Ala Ser Ala Lys	Arg Val Pro Leu Pro	Pro Met Thr Ser Thr Asp		
65	70	75		80
Tyr Val Asn Thr Ile	Pro Thr Ser Met Glu	Pro Asp Phe Pro Gly Asp		
	85	90		95
Glu Glu Met Glu Lys	Arg Tyr Arg Arg Trp	Met Arg Trp Asn Ala Ala		
	100	105		110
Ile Met Val His Arg	Ala Gln Arg Pro Gly	Ile Gly Val Gly Gly His		
	115	120		125
Ile Ser Thr Tyr Ala	Gly Ala Ala Pro Leu	Tyr Glu Val Gly Phe Asn		
	130	135		140
His Phe Phe Arg Gly	Lys Asp His Pro Gly	Gly Gly Asp Gln Val Phe		
	145	150		155
Phe Gln Gly His Ala	Ser Pro Gly Met Tyr	Ala Arg Ala Phe Leu Glu		
	165	170		175
Gly Arg Leu Thr Glu	Ser Asp Leu Asp Ser	Phe Arg Gln Glu Val Ser		
	180	185		190
Tyr Glu Gly Gly Gly	Ile Pro Ser Tyr Pro	His Pro His Gly Met Pro		
	195	200		205
Asp Phe Trp Glu Phe	Pro Thr Val Ser Met	Gly Leu Gly Pro Met Asp		
	210	215		220
Ala Ile Tyr Gln Ala	Arg Phe Asn Arg Tyr	Leu His Asn Arg Gly Ile		
	225	230		235
Lys Asp Thr Ser Glu	Gln His Val Trp Ala	Phe Leu Gly Asp Gly Glu		
	245	250		255
Met Asp Glu Pro Glu	Ser Arg Gly Leu Ile	His Gln Ala Ala Leu Asn		
	260	265		270
Asn Leu Asp Asn Leu	Thr Phe Val Ile Asn	Cys Asn Leu Gln Arg Leu		
	275	280		285
Asp Gly Pro Val Arg	Gly Asn Thr Lys Ile	Ile Gln Glu Leu Glu Ser		
	290	295		300
Phe Phe Arg Gly Ala	Gly Trp Ser Val Ile	Lys Val Ile Trp Gly Arg		
	305	310		315
Glu Trp Asp Glu Leu	Leu Glu Lys Asp Gln	Asp Gly Ala Leu Val Glu		
	325	330		335
Val Met Asn Asn Thr	Ser Asp Gly Asp Tyr	Gln Thr Phe Lys Ala Asn		
	340	345		350
Asp Gly Ala Tyr Val	Arg Glu His Phe Phe	Gly Arg Asp Pro Arg Thr		
	355	360		365
Leu Lys Leu Val Glu	Asp Met Thr Asp Glu	Glu Ile Trp Lys Leu Pro		
	370	375		380
Arg Gly Gly His Asp	Tyr Arg Lys Val Tyr	Ala Ala Tyr Lys Arg Ala		
	385	390		395
				400

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Leu Glu Thr Lys Asp Arg Pro Thr Val Ile Leu Ala His Thr Ile Lys  
 405 410 415  
 Gly Tyr Gly Leu Gly His Asn Phe Glu Gly Arg Asn Ala Thr His Gln  
 420 425 430  
 Met Lys Lys Leu Thr Leu Asp Asp Leu Lys Leu Phe Arg Asp Lys Gln  
 435 440 445  
 Gly Leu Pro Ile Thr Asp Glu Glu Leu Glu Lys Asp Pro Tyr Leu Pro  
 450 455 460  
 Pro Tyr Tyr His Pro Gly Glu Asp Ala Pro Glu Ile Lys Tyr Met Lys  
 465 470 475 480  
 Glu Arg Arg Gln Ala Leu Gly Gly Phe Leu Pro Glu Arg Arg Glu Lys  
 485 490 495  
 Tyr Glu Pro Leu Gln Val Pro Pro Leu Asp Lys Leu Arg Ser Val Arg  
 500 505 510  
 Lys Gly Ser Gly Lys Gln Gln Val Ala Thr Thr Met Ala Thr Val Arg  
 515 520 525  
 Thr Phe Lys Glu Leu Met Arg Asp Lys Asn Leu Ala Asp Arg Leu Val  
 530 535 540  
 Pro Ile Ile Pro Asp Glu Ala Arg Thr Phe Gly Leu Asp Ser Trp Phe  
 545 550 555 560  
 Pro Thr Leu Lys Ile Tyr Asn Pro His Gly Gln Asn Tyr Val Pro Val  
 565 570 575  
 Asp His Asp Leu Met Leu Ser Tyr Arg Glu Ala Lys Asp Gly Gln Ile  
 580 585 590  
 Leu His Glu Gly Ile Asn Glu Ala Gly Ser Val Ala Ser Phe Ile Ala  
 595 600 605  
 Ala Gly Thr Ser Tyr Ala Thr His Gly Glu Ala Met Ile Pro Leu Tyr  
 610 615 620  
 Ile Phe Tyr Ser Met Phe Gly Phe Gln Arg Thr Gly Asp Gly Ile Trp  
 625 630 635 640  
 Ala Ala Ala Asp Gln Met Thr Arg Gly Phe Leu Leu Gly Ala Thr Ala  
 645 650 655  
 Gly Arg Thr Thr Leu Thr Gly Glu Gly Leu Gln His Met Asp Gly His  
 660 665 670  
 Ser Pro Ile Leu Ala Ser Thr Asn Pro Gly Val Glu Thr Tyr Asp Pro  
 675 680 685  
 Ala Phe Ser Tyr Glu Ile Ala His Leu Val His Arg Gly Ile Asp Arg  
 690 695 700  
 Met Tyr Gly Pro Gly Lys Gly Glu Asn Val Ile Tyr Tyr Leu Thr Ile  
 705 710 715 720  
 Tyr Asn Glu Pro Thr Pro Gln Pro Ala Glu Pro Glu Asp Leu Asp Val  
 725 730 735  
 Glu Gly Leu His Lys Gly Ile Tyr Leu Tyr Asp Lys Ala Ala Glu Gly

gtcctttt	caaattctgc	aaagtgggta	gaggtcagat	gtcagcaggi	cggltccgatt	60
tcgttaggaa	agltggagccg	tggggggcaa	caltaaccit	ccccctggga	tgtagctaaa	120
cggcaalggg	ggtctcgggc	ggggggcatl	cttttcacgg	caaggttggtg	aaatlccgca	180
ggltcaciccc	cggccggcgg	tagagaacgg	agcgaaaacg	gaaagcaata	cgtggttttc	240
cggactlggcc	gttacgaigt	tcigaagagt	gactgccatc	acceaacagg	ctggctccicg	300
tcgaaaggaa	caaaaaact	gtg gti aca	aca aca ccc	tcc acg ctg	cgc gcg	351
		Val Val Thr	Thr Thr Pro	Ser Thr Leu	Pro Ala	
		1	5	10		
ttc aaa aag atc	cig gig gcc	aac cga ggi	gaa atc gcg	gtg cga gca		399

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Phe	Lys	Lys	Ile	Leu	Val	Ala	Asn	Arg	Gly	Glu	Ile	Ala	Val	Arg	Ala	
			15					20					25			
ttc	cgc	gcc	gcc	tac	gag	acc	ggg	gcc	gca	acc	gtg	gcc	atc	tac	ccc	447
Phe	Arg	Ala	Ala	Tyr	Glu	Thr	Gly	Ala	Ala	Thr	Val	Ala	Ile	Tyr	Pro	
		30					35					40				
cgg	gag	gac	cgt	ggc	tcc	ttc	cac	cgc	tcc	ttc	gcc	tcc	gag	gcg	gtg	495
Arg	Glu	Asp	Arg	Gly	Ser	Phe	His	Arg	Ser	Phe	Ala	Ser	Glu	Ala	Val	
	45					50				55						
agg	atc	gga	acc	gag	ggc	tca	ccc	gtc	aag	gcg	tac	ctc	gat	att	gat	543
Arg	Ile	Gly	Thr	Glu	Gly	Ser	Pro	Val	Lys	Ala	Tyr	Leu	Asp	Ile	Asp	
60					65				70							
gag	atc	atc	aac	gcc	gcc	aag	aag	gtg	aaa	gcg	gac	gcg	gtc	tac	ccg	591
Glu	Ile	Ile	Asn	Ala	Ala	Lys	Lys	Val	Lys	Ala	Asp	Ala	Val	Tyr	Pro	
			80					85					90			
ggg	tat	ggt	ttc	cit	tcg	gaa	aat	gcc	cag	ctc	gcg	cgt	gaa	tgc	gcg	639
Gly	Tyr	Gly	Phe	Leu	Ser	Glu	Asn	Ala	Gln	Leu	Ala	Arg	Glu	Cys	Ala	
			95				100					105				
gag	aac	ggc	att	acc	ttc	atc	ggt	ccc	acc	ccg	gag	gtg	ctc	gac	ctc	687
Glu	Asn	Gly	Ile	Thr	Phe	Ile	Gly	Pro	Thr	Pro	Glu	Val	Leu	Asp	Leu	
	110						115				120					
acg	ggc	gac	aag	tcc	aag	gct	gtg	tcc	gcc	gcg	aag	aag	gcc	ggg	ctg	735
Thr	Gly	Asp	Lys	Ser	Lys	Ala	Val	Ser	Ala	Ala	Lys	Lys	Ala	Gly	Leu	
	125					130					135					
ccg	gtg	ctg	gcg	gaa	tcc	acc	ccc	agc	acc	gac	atc	gat	gag	atc	gtc	783
Pro	Val	Leu	Ala	Glu	Ser	Thr	Pro	Ser	Thr	Asp	Ile	Asp	Glu	Ile	Val	
140					145					150					155	
aag	agt	gcc	gag	ggg	cag	acc	tac	ccg	atc	ttc	gtc	aag	gcc	gtc	gca	831
Lys	Ser	Ala	Glu	Gly	Gln	Thr	Tyr	Pro	Ile	Phe	Val	Lys	Ala	Val	Ala	
			160					165				170				
ggt	ggt	ggc	ggg	cgt	ggt	atg	cgg	ttc	gtc	gag	aag	ccc	gag	gac	ctg	879
Gly	Gly	Gly	Gly	Arg	Gly	Met	Arg	Phe	Val	Glu	Lys	Pro	Glu	Asp	Leu	
			175				180					185				
cgt	gag	ctg	gcc	agg	gag	gcc	tcc	cgc	gag	gcg	gag	gcc	gct	ttc	ggt	927
Arg	Glu	Leu	Ala	Arg	Glu	Ala	Ser	Arg	Glu	Ala	Glu	Ala	Ala	Phe	Gly	
	190						195				200					
gac	gga	tcc	gtc	tac	gtc	gaa	cgg	gcc	gtg	atc	aaa	ccc	cag	cac	atc	975
Asp	Gly	Ser	Val	Tyr	Val	Glu	Arg	Ala	Val	Ile	Lys	Pro	Gln	His	Ile	
	205					210				215						
gag	gtg	cag	atc	ctc	ggt	gat	cac	acc	ggc	gat	gtc	atc	cac	ctg	tat	1023
Glu	Val	Gln	Ile	Leu	Gly	Asp	His	Thr	Gly	Asp	Val	Ile	His	Leu	Tyr	
220					225					230					235	
gaa	cgc	gac	tgt	tcc	ctg	cag	cgc	cgc	cac	cag	aag	gtc	gtg	gag	atc	1071
Glu	Arg	Asp	Cys	Ser	Leu	Gln	Arg	Arg	His	Gln	Lys	Val	Val	Glu	Ile	

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240										245					250					
gca	cct	gcc	cag	cac	ctc	gac	ccg	gag	ctg	cgc	gac	cgc	atc	tgt	gcc	1119				
Ala	Pro	Ala	Gln	His	Leu	Asp	Pro	Glu	Leu	Arg	Asp	Arg	Ile	Cys	Ala					
255										260					265					
gat	gcc	gtg	aag	ttc	tgc	aaa	icc	atc	gga	tac	cag	ggc	gcc	ggc	acc	1167				
Asp	Ala	Val	Lys	Phe	Cys	Lys	Ser	Ile	Gly	Tyr	Gln	Gly	Ala	Gly	Thr					
270										275					280					
gtg	gag	ttc	ctc	gtc	gac	gag	gcg	ggc	aac	cac	gtc	ttc	att	gag	atg	1215				
Val	Glu	Phe	Leu	Val	Asp	Glu	Ala	Gly	Asn	His	Val	Phe	Ile	Glu	Met					
285										290					295					
aac	ccc	cgc	atc	cag	gtg	gaa	cac	acc	gtg	acc	gag	gag	gtc	acc	tcc	1263				
Asn	Pro	Arg	Ile	Gln	Val	Glu	His	Thr	Val	Thr	Glu	Glu	Val	Thr	Ser					
300										305					310					
gtc	gac	ctg	gtc	aag	gcg	cag	atg	cac	ctg	gcc	gcc	ggt	gcc	acc	ctg	1311				
Val	Asp	Leu	Val	Lys	Ala	Gln	Met	His	Leu	Ala	Ala	Gly	Ala	Thr	Leu					
320										325					330					
aag	gaa	ctg	ggc	ctg	acc	cag	gac	aag	atc	acc	acc	cac	ggt	gcc	gcc	1359				
Lys	Glu	Leu	Gly	Leu	Thr	Gln	Asp	Lys	Ile	Thr	Thr	His	Gly	Ala	Ala					
335										340					345					
ctg	cag	tgc	cgc	atc	acc	acg	gag	gac	ccg	tcc	aac	aac	ttc	cgg	ccc	1407				
Leu	Gln	Cys	Arg	Ile	Thr	Thr	Glu	Asp	Pro	Ser	Asn	Asn	Phe	Arg	Pro					
350										355					360					
gac	acc	ggt	gtg	atc	acc	gcc	tac	cgc	tcc	ccg	ggt	ggt	gcg	ggt	gtg	1455				
Asp	Thr	Gly	Val	Ile	Thr	Ala	Tyr	Arg	Ser	Pro	Gly	Gly	Ala	Gly	Val					
365										370					375					
cgt	ctc	gac	ggc	gca	gcc	cag	ctc	ggc	ggc	gag	atc	acc	gca	cat	ttc	1503				
Arg	Leu	Asp	Gly	Ala	Ala	Gln	Leu	Gly	Gly	Glu	Ile	Thr	Ala	His	Phe					
380										385					390					
gat	tcc	atg	ctg	gtc	aag	atg	acc	tgc	cgc	ggt	tcc	gat	ttc	gag	acc	1551				
Asp	Ser	Met	Leu	Val	Lys	Met	Thr	Cys	Arg	Gly	Ser	Asp	Phe	Glu	Thr					
400										405					410					
gcc	gtg	tcc	cga	gcc	cag	cgc	gcc	ctg	gcg	gag	ttc	aac	gtc	tcc	ggc	1599				
Ala	Val	Ser	Arg	Ala	Gln	Arg	Ala	Leu	Ala	Glu	Phe	Asn	Val	Ser	Gly					
415										420					425					
gtg	gcc	acc	aac	atc	ggc	ttc	ctg	cgt	gcg	ctg	ctg	cgc	gag	gaa	gac	1647				
Val	Ala	Thr	Asn	Ile	Gly	Phe	Leu	Arg	Ala	Leu	Leu	Arg	Glu	Glu	Asp					
430										435					440					
ttc	acc	aag	agg	cgc	atc	gac	acc	ggc	ttc	atc	ggc	tcc	cac	cag	cac	1695				
Phe	Thr	Lys	Arg	Arg	Ile	Asp	Thr	Gly	Phe	Ile	Gly	Ser	His	Gln	His					
445										450					455					
ctg	ctc	cag	gcc	cca	ccg	gcc	gac	gat	gag	cag	ggg	cgg	atc	ctg	gaa	1743				
Leu	Leu	Gln	Ala	Pro	Pro	Ala	Asp	Asp	Glu	Gln	Gly	Arg	Ile	Leu	Glu					
460										465					470					
															475					



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tac ctg gcg gat gtc acc gtg aac aaa ccc cac ggt gaa cgc ccc gag	1791
Tyr Leu Ala Asp Val Thr Val Asn Lys Pro His Gly Glu Arg Pro Glu	
480 485 490	
aca gcc cgt ccg ata gag aag ctg ccc gag gtg gag aac atc ccg ctg	1839
Thr Ala Arg Pro Ile Glu Lys Leu Pro Glu Val Glu Asn Ile Pro Leu	
495 500 505	
cca cgc ggc tcc cgc gac cgc ctg aag cag ctg ggc ccg gag ggt ttc	1887
Pro Arg Gly Ser Arg Asp Arg Leu Lys Gln Leu Gly Pro Glu Gly Phe	
510 515 520	
gcc cgc gat ctg cgc gaa cag gat gcc ctg gcc gtc acc gac acc acc	1935
Ala Arg Asp Leu Arg Glu Gln Asp Ala Leu Ala Val Thr Asp Thr Thr	
525 530 535	
ttc cgc gat gcc cac cag tcc ctg ctg gcc acc cgc gtg cgc tcc ttc	1983
Phe Arg Asp Ala His Gln Ser Leu Leu Ala Thr Arg Val Arg Ser Phe	
540 545 550 555	
gcg ctg acc ccg gcg gcg cgc gcc gtc gca aag ctg acc ccc gag ctg	2031
Ala Leu Thr Pro Ala Ala Arg Ala Val Ala Lys Leu Thr Pro Glu Leu	
560 565 570	
ctg tgc gtg gag gcc tgg ggc ggt gcc acc tac gac gtg gcc atg cgc	2079
Leu Ser Val Glu Ala Trp Gly Gly Ala Thr Tyr Asp Val Ala Met Arg	
575 580 585	
ttc ctg ttc gag gat ccg tgg gca cgc ctg gat gag ctg cgt gag gcg	2127
Phe Leu Phe Glu Asp Pro Trp Ala Arg Leu Asp Glu Leu Arg Glu Ala	
590 595 600	
atg ccg aat gtg aac atc cag atg ctg ctg cgt ggt cgc aac acc gtc	2175
Met Pro Asn Val Asn Ile Gln Met Leu Leu Arg Gly Arg Asn Thr Val	
605 610 615	
ggg tac acc ccg tac ccc gat tgc gtg tgc cgc gcg ttt gtg cag gag	2223
Gly Tyr Thr Pro Tyr Pro Asp Ser Val Cys Arg Ala Phe Val Gln Glu	
620 625 630 635	
gcc gcc aag tcc ggt gtg gac atc ttc cgc atc ttc gac gcg ctg aac	2271
Ala Ala Lys Ser Gly Val Asp Ile Phe Arg Ile Phe Asp Ala Leu Asn	
640 645 650	
gac atc tcc cag atg cgc ccg gcc atc gac gcc gtc ctg gag acc gcc	2319
Asp Ile Ser Gln Met Arg Pro Ala Ile Asp Ala Val Leu Glu Thr Gly	
655 660 665	
acc agt gtt gcc gag gtc gcc atg gcg tac tcc ggt gac ctg tcc aat	2367
Thr Ser Val Ala Glu Val Ala Met Ala Tyr Ser Gly Asp Leu Ser Asn	
670 675 680	
ccg ggg gag aag ctg tac acc ctg gac tac tac ctg aac ctg gcc gag	2415
Pro Gly Glu Lys Leu Tyr Thr Leu Asp Tyr Tyr Leu Asn Leu Ala Glu	
685 690 695	
cag atc gtc gac tcc ggt gca cac atc ctg gcc atc aag gac atg gcc	2463

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Gln	Ile	Val	Asp	Ser	Gly	Ala	His	Ile	Leu	Ala	Ile	Lys	Asp	Met	Ala		
700					705					710					715		
ggc	ctg	ctg	cgc	cgc	gcc	gcg	gcg	ccc	aaa	ctg	gtc	acc	gcc	ctg	cgc	2511	
Gly	Leu	Leu	Arg	Arg	Ala	Ala	Ala	Pro	Lys	Leu	Val	Thr	Ala	Leu	Arg		
				720					725					730			
cgt	gaa	ttc	gac	ctg	ccc	gtg	cat	gtc	cac	acc	cac	gac	acc	gcc	ggc	2559	
Arg	Glu	Phe	Asp	Leu	Pro	Val	His	Val	His	Thr	His	Asp	Thr	Ala	Gly		
			735					740					745				
ggt	cag	ctg	gcc	acc	tac	ctg	gcc	gcc	gcc	aac	gcc	ggg	gcc	gat	gcc	2607	
Gly	Gln	Leu	Ala	Thr	Tyr	Leu	Ala	Ala	Ala	Asn	Ala	Gly	Ala	Asp	Ala		
		750						755				760					
gtc	gac	gcc	gcc	tcc	gca	ccc	ctg	tcc	ggt	acc	acc	tcc	cag	ccg	tcc	2655	
Val	Asp	Ala	Ala	Ser	Ala	Pro	Leu	Ser	Gly	Thr	Thr	Ser	Gln	Pro	Ser		
		765				770						775					
atg	tcc	gct	ctg	gtt	gcc	gcg	ttt	gcg	cac	acc	cga	cgc	gac	acc	ggc	2703	
Met	Ser	Ala	Leu	Val	Ala	Ala	Phe	Ala	His	Thr	Arg	Arg	Asp	Thr	Gly		
780					785					790					795		
ctc	aac	ctg	cag	gcc	gtc	tcc	gac	ctg	gaa	ccg	tac	tgg	gag	gcg	gtc	2751	
Leu	Asn	Leu	Gln	Ala	Val	Ser	Asp	Leu	Glu	Pro	Tyr	Trp	Glu	Ala	Val		
				800					805				810				
cgc	gga	ctg	tac	ctg	ccg	ttt	gaa	tcc	ggc	acc	ccg	ggc	ccg	acc	gga	2799	
Arg	Gly	Leu	Tyr	Leu	Pro	Phe	Glu	Ser	Gly	Thr	Pro	Gly	Pro	Thr	Gly		
			815					820				825					
cgc	gtt	tac	cgc	cac	gag	atc	ccc	ggc	ggt	cag	ctg	tcc	aac	ctg	cgt	2847	
Arg	Val	Tyr	Arg	His	Glu	Ile	Pro	Gly	Gly	Gln	Leu	Ser	Asn	Leu	Arg		
		830					835					840					
gcc	cag	gcc	gtt	gca	ctg	ggt	ctg	gcc	gac	cgc	ttc	gag	ctc	atc	gag	2895	
Ala	Gln	Ala	Val	Ala	Leu	Gly	Leu	Ala	Asp	Arg	Phe	Glu	Leu	Ile	Glu		
		845				850				855							
gac	tac	tac	gcg	gcc	gtc	aac	gag	atg	ctg	ggt	cgt	ccg	acc	aag	gtc	2943	
Asp	Tyr	Tyr	Ala	Ala	Val	Asn	Glu	Met	Leu	Gly	Arg	Pro	Thr	Lys	Val		
860					865				870					875			
acc	ccg	tcc	tcc	aag	gtt	gtc	ggt	gac	ctc	gca	ctg	cac	ctc	gtc	ggt	2991	
Thr	Pro	Ser	Ser	Lys	Val	Val	Gly	Asp	Leu	Ala	Leu	His	Leu	Val	Gly		
				880					885					890			
gcc	ggt	gtg	agc	ccg	gag	gat	ttc	gcc	gcc	gat	ccg	cag	aag	tac	gac	3039	
Ala	Gly	Val	Ser	Pro	Glu	Asp	Phe	Ala	Ala	Asp	Pro	Gln	Lys	Tyr	Asp		
				895				900				905					
atc	ccc	gat	tcc	gtc	atc	gcc	ttc	ctc	cgc	ggc	gaa	ctg	ggt	acc	ccg	3087	
Ile	Pro	Asp	Ser	Val	Ile	Ala	Phe	Leu	Arg	Gly	Glu	Leu	Gly	Thr	Pro		
		910				915					920						
ccc	ggt	ggc	tgg	ccc	gaa	ccg	ctg	cgc	acc	cgt	gca	ctc	gag	ggt	cgc	3135	
Pro	Gly	Gly	Trp	Pro	Glu	Pro	Leu	Arg	Thr	Arg	Ala	Leu	Glu	Gly	Arg		

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925	930	935	
icc cag ggt aag gcc ccg ctg gcg gag atc ccc gcc gag gag cag gcc			3183
Ser Gln Gly Lys Ala Pro Leu Ala Glu Ile Pro Ala Glu Glu Gln Ala			
940	945	950	955
cac ctg gat tcc gat gat tcc gcg gag cgt cgc ggc acc ctg aac cgc			3231
His Leu Asp Ser Asp Asp Ser Ala Glu Arg Arg Gly Thr Leu Asn Arg			
960	965	970	
ctg ctg ttc ccg aag ccg acc gag gag ttc ctt gag cac cgt cgc cgc			3279
Leu Leu Phe Pro Lys Pro Thr Glu Glu Phe Leu Glu His Arg Arg Arg			
975	980	985	
ttc ggc aac acc tcc gcc ctg gat gac cgc gag ttc ttc tac ggc ttg			3327
Phe Gly Asn Thr Ser Ala Leu Asp Asp Arg Glu Phe Phe Tyr Gly Leu			
990	995	1000	
aag gag gga cgt gag gag ctg atc cga ctg acc ggt gtg tcc acc ccg			3375
Lys Glu Gly Arg Glu Glu Leu Ile Arg Leu Thr Gly Val Ser Thr Pro			
1005	1010	1015	
atg gtg gtc cgc ctg gat gcg gtg tcc gaa ccg gat gac aaa ggc atg			3423
Met Val Val Arg Leu Asp Ala Val Ser Glu Pro Asp Asp Lys Gly Met			
1020	1025	1030	1035
cgc aac gtg gtg gtc aac gtc aac ggc cag atc cgc ccg atc aag gtg			3471
Arg Asn Val Val Val Asn Val Asn Gly Gln Ile Arg Pro Ile Lys Val			
1040	1045	1050	
cgc gac cgt tcc gtg gag tcc gtc acc gcc acc gcg gag aag gcc gat			3519
Arg Asp Arg Ser Val Glu Ser Val Thr Ala Thr Ala Glu Lys Ala Asp			
1055	1060	1065	
gcc acc aac aag ggc cat gtc gcc gca cca ttc gcc ggt gtg gtc acc			3567
Ala Thr Asn Lys Gly His Val Ala Ala Pro Phe Ala Gly Val Val Thr			
1070	1075	1080	
gtg acc gtc gcc gag ggt gat gag atc aag gct ggc gac gcc gtg gcc			3615
Val Thr Val Ala Glu Gly Asp Glu Ile Lys Ala Gly Asp Ala Val Ala			
1085	1090	1095	
atc att gag gcc atg aag atg gag gcc acc atc acc gcg cct gtc gac			3663
Ile Ile Glu Ala Met Lys Met Glu Ala Thr Ile Thr Ala Pro Val Asp			
1100	1105	1110	1115
ggt gtc atc gac cgc gtc gtc gtc ccc gcc gcc acc aag gtc gag ggc			3711
Gly Val Ile Asp Arg Val Val Val Pro Ala Ala Thr Lys Val Glu Gly			
1120	1125	1130	
ggc gac ctg atc gtg gtc gtg tcc tagcgactga gagccacaac ccgtcccg			3765
Gly Asp Leu Ile Val Val Val Ser			
1135			
tgccttgta tcaacctccc cctgatgatg ttctcagggg gaggcctctac gtacctcacc			3825
gtgacgggic atgtatctg tctgtctgga gagaatgctc caggtaggaa cgccaaccac			3885
cccactccgt gatgtccgt gctgatccca ggcaggccgg ttggaaagaa aaaccagtga			3945

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tggaacggcc atcggacagc gagacggaac caagcgatc cggctccggt agagcggiga 4005  
ggagcctg 4013

&lt;210&gt; 24

&lt;211&gt; 1139

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 24

Val	Val	Thr	Thr	Thr	Pro	Ser	Thr	Leu	Pro	Ala	Phe	Lys	Lys	Ile	Leu
1				5					10					15	
Val	Ala	Asn	Arg	Gly	Glu	Ile	Ala	Val	Arg	Ala	Phe	Arg	Ala	Ala	Tyr
		20						25				30			
Glu	Thr	Gly	Ala	Ala	Thr	Val	Ala	Ile	Tyr	Pro	Arg	Glu	Asp	Arg	Gly
		35					40					45			
Ser	Phe	His	Arg	Ser	Phe	Ala	Ser	Glu	Ala	Val	Arg	Ile	Gly	Thr	Glu
	50					55					60				
Gly	Ser	Pro	Val	Lys	Ala	Tyr	Leu	Asp	Ile	Asp	Glu	Ile	Ile	Asn	Ala
65					70				75					80	
Ala	Lys	Lys	Val	Lys	Ala	Asp	Ala	Val	Tyr	Pro	Gly	Tyr	Gly	Phe	Leu
			85					90						95	
Ser	Glu	Asn	Ala	Gln	Leu	Ala	Arg	Glu	Cys	Ala	Glu	Asn	Gly	Ile	Thr
		100						105				110			
Phe	Ile	Gly	Pro	Thr	Pro	Glu	Val	Leu	Asp	Leu	Thr	Gly	Asp	Lys	Ser
	115					120					125				
Lys	Ala	Val	Ser	Ala	Ala	Lys	Lys	Ala	Gly	Leu	Pro	Val	Leu	Ala	Glu
	130					135					140				
Ser	Thr	Pro	Ser	Thr	Asp	Ile	Asp	Glu	Ile	Val	Lys	Ser	Ala	Glu	Gly
145					150				155					160	
Gln	Thr	Tyr	Pro	Ile	Phe	Val	Lys	Ala	Val	Ala	Gly	Gly	Gly	Gly	Arg
			165					170						175	
Gly	Met	Arg	Phe	Val	Glu	Lys	Pro	Glu	Asp	Leu	Arg	Glu	Leu	Ala	Arg
		180						185				190			
Glu	Ala	Ser	Arg	Glu	Ala	Glu	Ala	Ala	Phe	Gly	Asp	Gly	Ser	Val	Tyr
		195				200					205				
Val	Glu	Arg	Ala	Val	Ile	Lys	Pro	Gln	His	Ile	Glu	Val	Gln	Ile	Leu
	210					215					220				
Gly	Asp	His	Thr	Gly	Asp	Val	Ile	His	Leu	Tyr	Glu	Arg	Asp	Cys	Ser
225					230				235					240	
Leu	Gln	Arg	Arg	His	Gln	Lys	Val	Val	Glu	Ile	Ala	Pro	Ala	Gln	His
			245						250					255	
Leu	Asp	Pro	Glu	Leu	Arg	Asp	Arg	Ile	Cys	Ala	Asp	Ala	Val	Lys	Phe
		260						265				270			
Cys	Lys	Ser	Ile	Gly	Tyr	Gln	Gly	Ala	Gly	Thr	Val	Glu	Phe	Leu	Val

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275	280	285
Asp Glu Ala Gly Asn His Val Phe Ile Glu Met Asn Pro Arg Ile Gln		
290	295	300
Val Glu His Thr Val Thr Glu Glu Val Thr Ser Val Asp Leu Val Lys		
305	310	315
Ala Gln Met His Leu Ala Ala Gly Ala Thr Leu Lys Glu Leu Gly Leu		
	325	330
Thr Gln Asp Lys Ile Thr Thr His Gly Ala Ala Leu Gln Cys Arg Ile		
	340	345
Thr Thr Glu Asp Pro Ser Asn Asn Phe Arg Pro Asp Thr Gly Val Ile		
	355	360
Thr Ala Tyr Arg Ser Pro Gly Glu Ala Gly Val Arg Leu Asp Gly Ala		
	370	375
Ala Gln Leu Gly Gly Glu Ile Thr Ala His Phe Asp Ser Met Leu Val		
385	390	395
Lys Met Thr Cys Arg Gly Ser Asp Phe Glu Thr Ala Val Ser Arg Ala		
	405	410
Gln Arg Ala Leu Ala Glu Phe Asn Val Ser Gly Val Ala Thr Asn Ile		
	420	425
Gly Phe Leu Arg Ala Leu Leu Arg Glu Glu Asp Phe Thr Lys Arg Arg		
	435	440
Ile Asp Thr Gly Phe Ile Gly Ser His Gln His Leu Leu Gln Ala Pro		
	450	455
Pro Ala Asp Asp Glu Gln Gly Arg Ile Leu Glu Tyr Leu Ala Asp Val		
465	470	475
Thr Val Asn Lys Pro His Gly Glu Arg Pro Glu Thr Ala Arg Pro Ile		
	485	490
Glu Lys Leu Pro Glu Val Glu Asn Ile Pro Leu Pro Arg Gly Ser Arg		
	500	505
Asp Arg Leu Lys Gln Leu Gly Pro Glu Gly Phe Ala Arg Asp Leu Arg		
	515	520
Glu Gln Asp Ala Leu Ala Val Thr Asp Thr Thr Phe Arg Asp Ala His		
	530	535
Gln Ser Leu Leu Ala Thr Arg Val Arg Ser Phe Ala Leu Thr Pro Ala		
545	550	555
Ala Arg Ala Val Ala Lys Leu Thr Pro Glu Leu Leu Ser Val Glu Ala		
	565	570
Trp Gly Gly Ala Thr Tyr Asp Val Ala Met Arg Phe Leu Phe Glu Asp		
	580	585
Pro Trp Ala Arg Leu Asp Glu Leu Arg Glu Ala Met Pro Asn Val Asn		
	595	600
Ile Gln Met Leu Leu Arg Gly Arg Asn Thr Val Gly Tyr Thr Pro Tyr		
610	615	620

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Pro Asp Ser Val Cys Arg Ala Phe Val Gln Glu Ala Ala Lys Ser Gly  
 625 630 635 640  
 Val Asp Ile Phe Arg Ile Phe Asp Ala Leu Asn Asp Ile Ser Gln Met  
 645 650 655  
 Arg Pro Ala Ile Asp Ala Val Leu Glu Thr Gly Thr Ser Val Ala Glu  
 660 665 670  
 Val Ala Met Ala Tyr Ser Gly Asp Leu Ser Asn Pro Gly Glu Lys Leu  
 675 680 685  
 Tyr Thr Leu Asp Tyr Tyr Leu Asn Leu Ala Glu Gln Ile Val Asp Ser  
 690 695 700  
 Gly Ala His Ile Leu Ala Ile Lys Asp Met Ala Gly Leu Leu Arg Arg  
 705 710 715 720  
 Ala Ala Ala Pro Lys Leu Val Thr Ala Leu Arg Arg Glu Phe Asp Leu  
 725 730 735  
 Pro Val His Val His Thr His Asp Thr Ala Gly Gly Gln Leu Ala Thr  
 740 745 750  
 Tyr Leu Ala Ala Ala Asn Ala Gly Ala Asp Ala Val Asp Ala Ala Ser  
 755 760 765  
 Ala Pro Leu Ser Gly Thr Thr Ser Gln Pro Ser Met Ser Ala Leu Val  
 770 775 780  
 Ala Ala Phe Ala His Thr Arg Arg Asp Thr Gly Leu Asn Leu Gln Ala  
 785 790 795 800  
 Val Ser Asp Leu Glu Pro Tyr Trp Glu Ala Val Arg Gly Leu Tyr Leu  
 805 810 815  
 Pro Phe Glu Ser Gly Thr Pro Gly Pro Thr Gly Arg Val Tyr Arg His  
 820 825 830  
 Glu Ile Pro Gly Gly Gln Leu Ser Asn Leu Arg Ala Gln Ala Val Ala  
 835 840 845  
 Leu Gly Leu Ala Asp Arg Phe Glu Leu Ile Glu Asp Tyr Tyr Ala Ala  
 850 855 860  
 Val Asn Glu Met Leu Gly Arg Pro Thr Lys Val Thr Pro Ser Ser Lys  
 865 870 875 880  
 Val Val Gly Asp Leu Ala Leu His Leu Val Gly Ala Gly Val Ser Pro  
 885 890 895  
 Glu Asp Phe Ala Ala Asp Pro Gln Lys Tyr Asp Ile Pro Asp Ser Val  
 900 905 910  
 Ile Ala Phe Leu Arg Gly Glu Leu Gly Thr Pro Pro Gly Gly Trp Pro  
 915 920 925  
 Glu Pro Leu Arg Thr Arg Ala Leu Glu Gly Arg Ser Gln Gly Lys Ala  
 930 935 940  
 Pro Leu Ala Glu Ile Pro Ala Glu Glu Gln Ala His Leu Asp Ser Asp  
 945 950 955 960  
 Asp Ser Ala Glu Arg Arg Gly Thr Leu Asn Arg Leu Leu Phe Pro Lys

<400> 25  
galcaaccia agccaggaga atccggcggg cggtttctac ttctacagga gctgaacccc 60  
acc gtg aat gaa ctt ctc cgt gac gal atc cgt tat ctc ggc cgg atc 108  
Val Asn Glu Leu Leu Arg Asp Asp Ile Arg Tyr Leu Gly Arg Ile  
1 5 10 15  
ctg ggc gag gtg atc tcc gag cag gag ggc cac cat gtc ttc gaa ctg 156  
Leu Gly Glu Val Ile Ser Glu Gln Glu Gly His His Val Phe Glu Leu  
20 25 30  
gtt gaa cgc gcc cgc cgg acc tcc ttc gac atc gcc aag gga cgc gcg 204  
Val Glu Arg Ala Arg Arg Thr Ser Phe Asp Ile Ala Lys Gly Arg Ala

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35				40				45									
gag	atg	gac	agt	ctg	gtg	gag	gig	ttc	gct	ggc	atc	gac	ccg	gag	gac	252	
Glu	Met	Asp	Ser	Leu	Val	Glu	Val	Phe	Ala	Gly	Ile	Asp	Pro	Glu	Asp		
50				55				60									
gcc	acg	ccc	gtg	gcc	cga	gcc	ttc	acc	cat	ttc	gcc	ctg	ttg	gcc	aac	300	
Ala	Thr	Pro	Val	Ala	Arg	Ala	Phe	Thr	His	Phe	Ala	Leu	Leu	Ala	Asn		
65				70				75									
ctc	gcg	gag	gat	ttg	cat	gac	gca	gcc	cag	cgg	gaa	cag	gcc	ctg	aac	348	
Leu	Ala	Glu	Asp	Leu	His	Asp	Ala	Ala	Gln	Arg	Glu	Gln	Ala	Leu	Asn		
80				85				90				95					
tcg	ggt	gag	ccc	gcg	ccg	gac	agc	acc	ctc	gag	gcc	acc	tgg	gtg	aaa	396	
Ser	Gly	Glu	Pro	Ala	Pro	Asp	Ser	Thr	Leu	Glu	Ala	Thr	Trp	Val	Lys		
100				105				110									
ctg	gat	gat	gcc	ggg	gtg	ggc	agc	ggt	gag	gtc	gcc	gcg	gtg	atc	cgc	444	
Leu	Asp	Asp	Ala	Gly	Val	Gly	Ser	Gly	Glu	Val	Ala	Ala	Val	Ile	Arg		
115				120				125									
aat	gcg	ctc	gtc	gcc	ccg	gtg	ctc	acc	gcg	cac	ccg	acg	gaa	acc	cga	492	
Asn	Ala	Leu	Val	Ala	Pro	Val	Leu	Thr	Ala	His	Pro	Thr	Glu	Thr	Arg		
130				135				140									
cgt	cgt	acc	gtg	ttc	gac	gcg	cag	aag	cac	atc	acc	gcc	ctg	atg	gag	540	
Arg	Arg	Thr	Val	Phe	Asp	Ala	Gln	Lys	His	Ile	Thr	Ala	Leu	Met	Glu		
145				150				155									
gaa	cgc	cac	ctc	ctc	ctg	gcg	ctg	ccc	acg	cat	gcc	cgg	acc	cag	tcc	588	
Glu	Arg	His	Leu	Leu	Leu	Ala	Leu	Pro	Thr	His	Ala	Arg	Thr	Gln	Ser		
160				165				170				175					
aag	ctg	gat	gac	atc	gag	cgc	aac	atc	cgg	cga	cgg	atc	acg	atc	ctg	636	
Lys	Leu	Asp	Asp	Ile	Glu	Arg	Asn	Ile	Arg	Arg	Arg	Ile	Thr	Ile	Leu		
180				185				190									
tgg	cag	acg	gcc	ctc	atc	cgt	gtg	gcc	cgt	ccc	cgc	atc	gag	gat	gag	684	
Trp	Gln	Thr	Ala	Leu	Ile	Arg	Val	Ala	Arg	Pro	Arg	Ile	Glu	Asp	Glu		
195				200				205									
gtc	gag	ggt	gga	ctg	cgc	tac	tac	aag	ctc	agc	ctg	ttg	gcc	gag	atc	732	
Val	Glu	Val	Gly	Leu	Arg	Tyr	Tyr	Lys	Leu	Ser	Leu	Leu	Ala	Glu	Ile		
210				215				220									
ccc	cgc	atc	aat	cat	gat	gtg	acc	gtg	gaa	ctg	gcc	cgg	cgt	ttc	ggc	780	
Pro	Arg	Ile	Asn	His	Asp	Val	Thr	Val	Glu	Leu	Ala	Arg	Arg	Phe	Gly		
225				230				235									
ggg	gat	atc	ccc	acc	acg	gcg	atg	gtc	agg	ccg	gga	icc	tgg	atc	ggc	828	
Gly	Asp	Ile	Pro	Thr	Thr	Ala	Met	Val	Arg	Pro	Gly	Ser	Trp	Ile	Gly		
240				245				250				255					
ggg	gac	cat	gat	ggc	aac	ccc	ttc	gtc	acc	gcg	gag	act	gtc	acc	tac	876	
Gly	Asp	His	Asp	Gly	Asn	Pro	Phe	Val	Thr	Ala	Glu	Thr	Val	Thr	Tyr		
260				265				270									



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gcc acc cat cgg gcc gcg gag acc gtg ctc aag tac tac gtc aag caa	924
Ala Thr His Arg Ala Ala Glu Thr Val Leu Lys Tyr Tyr Val Lys Gln	
275 280 285	
ctg cac gcc ctg gaa cac gaa ctc agt ctc tcc gac cgg atg aac gtc	972
Leu His Ala Leu Glu His Glu Leu Ser Leu Ser Asp Arg Met Asn Val	
290 295 300	
atc agc gat gag ctg cgt gtg ctt gcc gat gcc ggc cag aat gac atg	1020
Ile Ser Asp Glu Leu Arg Val Leu Ala Asp Ala Gly Gln Asn Asp Met	
305 310 315	
ccc agc cgg gtt gat gaa ccc tac cgg cgg gcc atc cac ggc atg cgt	1068
Pro Ser Arg Val Asp Glu Pro Tyr Arg Arg Ala Ile His Gly Met Arg	
320 325 330 335	
ggc cgg atg ctg gcc acc acg gcc gcc ctg atc ggt gag gag gcg gtc	1116
Gly Arg Met Leu Ala Thr Thr Ala Ala Leu Ile Gly Glu Glu Ala Val	
340 345 350	
gag ggc acc tgg ttc aag acc ttc acg ccc tat acc gat acc cac gag	1164
Glu Gly Thr Trp Phe Lys Thr Phe Thr Pro Tyr Thr Asp Thr His Glu	
355 360 365	
ttc aaa cgc gac ctc gat atc gtg gat ggt tcc ctg aga atg tcc cgg	1212
Phe Lys Arg Asp Leu Asp Ile Val Asp Gly Ser Leu Arg Met Ser Arg	
370 375 380	
gat gac atc atc gcc gat gac cgt ctg gcc atg ctg cgc tcc gcc ctg	1260
Asp Asp Ile Ile Ala Asp Asp Arg Leu Ala Met Leu Arg Ser Ala Leu	
385 390 395	
gac agc ttc ggg ttc aac ctc tac tcc ctg gat ctg cgc cag aat tcc	1308
Asp Ser Phe Gly Phe Asn Leu Tyr Ser Leu Asp Leu Arg Gln Asn Ser	
400 405 410 415	
gac ggt ttc gag gat gtc ctc acc gaa ttg ttc gcc acc gcc cag acc	1356
Asp Gly Phe Glu Asp Val Leu Thr Glu Leu Phe Ala Thr Ala Gln Thr	
420 425 430	
gag aag aac tac cgc ggg ttg acg gag gcg gag aag ctg gac ctg ctg	1404
Glu Lys Asn Tyr Arg Gly Leu Thr Glu Ala Glu Lys Leu Asp Leu Leu	
435 440 445	
atc cgc gaa ctg agc aca ccc cgc ccg ctc atc ccg cac ggg gac ccg	1452
Ile Arg Glu Leu Ser Thr Pro Arg Pro Leu Ile Pro His Gly Asp Pro	
450 455 460	
gac tac tcc gag gcc acc aac cgt gaa ctg ggg att ttt tcc aag gcc	1500
Asp Tyr Ser Glu Ala Thr Asn Arg Glu Leu Gly Ile Phe Ser Lys Ala	
465 470 475	
gcg gag gcc gtg cgt aaa ttc ggt cct ctc atg gtg ccg cac tgc atc	1548
Ala Glu Ala Val Arg Lys Phe Gly Pro Leu Met Val Pro His Cys Ile	
480 485 490 495	
atc tcc atg gcc tct tcc gtc acg gac atc ctc gaa ccg atg gtg ctg	1596

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Ile	Ser	Met	Ala	Ser	Ser	Val	Thr	Asp	Ile	Leu	Glu	Pro	Met	Val	Leu		
				500					505					510			
ctc	aag	gag	ttc	ggi	ctg	atc	cgg	gcc	aac	ggg	aag	aac	ccg	acg	ggc	1644	
Leu	Lys	Glu	Phe	Gly	Leu	Ile	Arg	Ala	Asn	Gly	Lys	Asn	Pro	Thr	Gly		
			515					520					525				
agc	gtc	gac	gtg	atc	ccg	ctg	ttc	gag	acg	atc	gat	gac	ctc	cag	cgt	1692	
Ser	Val	Asp	Val	Ile	Pro	Leu	Phe	Glu	Thr	Ile	Asp	Asp	Leu	Gln	Arg		
		530					535				540						
ggc	gcg	ggc	atc	ctg	gag	gaa	tig	tgg	gac	atc	gac	ctc	tac	cgc	aat	1740	
Gly	Ala	Gly	Ile	Leu	Glu	Glu	Leu	Trp	Asp	Ile	Asp	Leu	Tyr	Arg	Asn		
	545					550					555						
tac	cit	gag	cag	cgg	gac	aac	gtc	cag	gag	gtc	atg	ctg	ggg	tat	tcc	1788	
Tyr	Leu	Glu	Gln	Arg	Asp	Asn	Val	Gln	Glu	Val	Met	Leu	Gly	Tyr	Ser		
560				565				570				575					
gac	tcc	aac	aag	gac	ggc	ggg	tac	ttc	gcc	gcc	aac	tgg	gcg	cit	tac	1836	
Asp	Ser	Asn	Lys	Asp	Gly	Gly	Tyr	Phe	Ala	Ala	Asn	Trp	Ala	Leu	Tyr		
			580					585				590					
gac	gcg	gag	tta	cgc	ctg	gtc	gaa	cta	tgc	cgg	ggc	cgt	aat	gtc	aag	1884	
Asp	Ala	Glu	Leu	Arg	Leu	Val	Glu	Leu	Cys	Arg	Gly	Arg	Asn	Val	Lys		
		595					600				605						
ctc	cgt	ctc	ttc	cac	ggt	cgt	ggt	ggc	acg	gtg	ggt	cgt	ggc	ggt	ggc	1932	
Leu	Arg	Leu	Phe	His	Gly	Arg	Gly	Gly	Thr	Val	Gly	Arg	Gly	Gly	Gly		
	610					615					620						
ccc	tcc	tat	gat	gcg	atc	ctg	gcc	cag	ccc	aag	ggc	gcg	gtc	cgg	ggt	1980	
Pro	Ser	Tyr	Asp	Ala	Ile	Leu	Ala	Gln	Pro	Lys	Gly	Ala	Val	Arg	Gly		
	625					630				635							
gcg	gtg	cgg	gtg	act	gaa	cag	ggc	gag	atc	atc	tcc	gcg	aag	tac	ggt	2028	
Ala	Val	Arg	Val	Thr	Glu	Gln	Gly	Glu	Ile	Ile	Ser	Ala	Lys	Tyr	Gly		
640					645			650				655					
aac	ccg	gat	acg	gca	cgc	cgc	aac	cit	gag	gcc	ctg	gtg	tcc	gcg	acg	2076	
Asn	Pro	Asp	Thr	Ala	Arg	Arg	Asn	Leu	Glu	Ala	Leu	Val	Ser	Ala	Thr		
			660				665				670						
ctg	gag	gca	tgc	cit	ctg	gat	gat	gtg	gaa	ctg	ccc	aat	cgg	gaa	cgc	2124	
Leu	Glu	Ala	Ser	Leu	Leu	Asp	Asp	Val	Glu	Leu	Pro	Asn	Arg	Glu	Arg		
		675					680				685						
gcg	cac	cag	atc	atg	ggg	gag	atc	tgc	gag	tig	agc	ttc	cgc	agg	tac	2172	
Ala	His	Gln	Ile	Met	Gly	Glu	Ile	Ser	Glu	Leu	Ser	Phe	Arg	Arg	Tyr		
		690				695					700						
tca	tca	ctg	gtc	cat	gag	gat	ccc	gga	ttc	atc	cag	tac	ttc	acc	cag	2220	
Ser	Ser	Leu	Val	His	Glu	Asp	Pro	Gly	Phe	Ile	Gln	Tyr	Phe	Thr	Gln		
	705					710					715						
tcc	acc	ccc	ctg	cag	gag	atc	gga	tcc	ctc	aac	atc	ggt	tcc	cga	ccc	2268	
Ser	Thr	Pro	Leu	Gln	Glu	Ile	Gly	Ser	Leu	Asn	Ile	Gly	Ser	Arg	Pro		

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720	725	730	735	
icc tca cgt aaa cag acc aac acg gtg gag gai ctg cgt gcc atc ccg				2316
Ser Ser Arg Lys Gln Thr Asn Thr Val Glu Asp Leu Arg Ala Ile Pro				
	740	745	750	
tgg gtg ctg agc igg tcc cag tcc cgt gtc atg ctg ccg ggc tgg ttc				2364
Trp Val Leu Ser Trp Ser Gln Ser Arg Val Met Leu Pro Gly Trp Phe				
	755	760	765	
ggt gtg ggt acc gca ctg cgt gag tgg atc ggt gag ggg gag ggg gct				2412
Gly Val Gly Thr Ala Leu Arg Glu Trp Ile Gly Glu Gly Glu Gly Ala				
	770	775	780	
gcg gag cgc atc gcg gag ctg cag gaa ctg aac cgg tgc tgg ccg ttc				2460
Ala Glu Arg Ile Ala Glu Leu Gln Glu Leu Asn Arg Cys Trp Pro Phe				
	785	790	795	
ttc acc tcg gtg ctg gac aac atg gcc cag gtg atg agc aag gcg gaa				2508
Phe Thr Ser Val Leu Asp Asn Met Ala Gln Val Met Ser Lys Ala Glu				
800	805	810	815	
ctg cgc ctg gcc agg ttg tac gcc gai ctg atc ccg gai cgc gag gtg				2556
Leu Arg Leu Ala Arg Leu Tyr Ala Asp Leu Ile Pro Asp Arg Glu Val				
	820	825	830	
gcg gac cgg atc tat gag acc atc ttc ggg gag tat ttc ctg acc aag				2604
Ala Asp Arg Ile Tyr Glu Thr Ile Phe Gly Glu Tyr Phe Leu Thr Lys				
	835	840	845	
gag atg ttc tgc acc atc acc ggt tcc cag gac ctg ctg gai gac aac				2652
Glu Met Phe Cys Thr Ile Thr Gly Ser Gln Asp Leu Leu Asp Asp Asn				
	850	855	860	
ccg gcg ctg gcg cga tcg gtg cgc agt cgg ttc ccg tac ctg ctg ccg				2700
Pro Ala Leu Ala Arg Ser Val Arg Ser Arg Phe Pro Tyr Leu Leu Pro				
	865	870	875	
ctc aat gtc atc cag gtg gag atg atg cgc cgg tac cgg tcc ggt gai				2748
Leu Asn Val Ile Gln Val Glu Met Met Arg Arg Tyr Arg Ser Gly Asp				
880	885	890	895	
gag ggc acg gct gtc cca cgt aat atc cgc ctg acc atg aat gga ttg				2796
Glu Gly Thr Ala Val Pro Arg Asn Ile Arg Leu Thr Met Asn Gly Leu				
	900	905	910	
icc acg gcc ctg cgc aac tcg ggt tagggcgcca gacgccccgg gaaccgcac				2850
Ser Thr Ala Leu Arg Asn Ser Gly				
	915			
ccgtgtata ctgtctaaag ttgcccgtg tcatccgggc gtgatggata gacaacttaa				2910
cggcaaagga ttctccccac atggcaciga cgtttcaaat cgtctcgtt ctgccagcg				2970
tgtctatgac ggcttctgtc ctgcctgaca agggtaaggc cggaggctg tcaagccctt				3030
tccgtggigg cgtccagttc aacctctccg gtccacggg gtgggagaag aacctggacc				3090
gcgtaccat cctgaccgca gtcctctggg tgaatcgtat tctcgcgtc aacctcatcc				3150
aggcgtactc ctgaccctg atctttcaag gccgtccctt cggggcagge cttttttgca				3210

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ttctccaggt gatgtccatc acccaccggt tttaaactat tgaccgatag aaacacctgc 3270  
actagggttat ctgttatgca atagaaaata gtgcat 3306

&lt;210&gt; 26

&lt;211&gt; 919

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 26

Val	Asn	Glu	Leu	Leu	Arg	Asp	Asp	Ile	Arg	Tyr	Leu	Gly	Arg	Ile	Leu
1				5				10						15	
Gly	Glu	Val	Ile	Ser	Glu	Gln	Glu	Gly	His	His	Val	Phe	Glu	Leu	Val
			20					25					30		
Glu	Arg	Ala	Arg	Arg	Thr	Ser	Phe	Asp	Ile	Ala	Lys	Gly	Arg	Ala	Glu
		35					40					45			
Met	Asp	Ser	Leu	Val	Glu	Val	Phe	Ala	Gly	Ile	Asp	Pro	Glu	Asp	Ala
	50					55					60				
Thr	Pro	Val	Ala	Arg	Ala	Phe	Thr	His	Phe	Ala	Leu	Leu	Ala	Asn	Leu
65				70				75						80	
Ala	Glu	Asp	Leu	His	Asp	Ala	Ala	Gln	Arg	Glu	Gln	Ala	Leu	Asn	Ser
			85					90						95	
Gly	Glu	Pro	Ala	Pro	Asp	Ser	Thr	Leu	Glu	Ala	Thr	Trp	Val	Lys	Leu
		100						105					110		
Asp	Asp	Ala	Gly	Val	Gly	Ser	Gly	Glu	Val	Ala	Ala	Val	Ile	Arg	Asn
	115					120						125			
Ala	Leu	Val	Ala	Pro	Val	Leu	Thr	Ala	His	Pro	Thr	Glu	Thr	Arg	Arg
	130					135					140				
Arg	Thr	Val	Phe	Asp	Ala	Gln	Lys	His	Ile	Thr	Ala	Leu	Met	Glu	Glu
145				150				155						160	
Arg	His	Leu	Leu	Leu	Ala	Leu	Pro	Thr	His	Ala	Arg	Thr	Gln	Ser	Lys
			165					170						175	
Leu	Asp	Asp	Ile	Glu	Arg	Asn	Ile	Arg	Arg	Arg	Ile	Thr	Ile	Leu	Trp
		180						185				190			
Gln	Thr	Ala	Leu	Ile	Arg	Val	Ala	Arg	Pro	Arg	Ile	Glu	Asp	Glu	Val
	195					200						205			
Glu	Val	Gly	Leu	Arg	Tyr	Tyr	Lys	Leu	Ser	Leu	Leu	Ala	Glu	Ile	Pro
	210				215					220					
Arg	Ile	Asn	His	Asp	Val	Thr	Val	Glu	Leu	Ala	Arg	Arg	Phe	Gly	Gly
225				230				235						240	
Asp	Ile	Pro	Thr	Thr	Ala	Met	Val	Arg	Pro	Gly	Ser	Trp	Ile	Gly	Gly
			245					250						255	
Asp	His	Asp	Gly	Asn	Pro	Phe	Val	Thr	Ala	Glu	Thr	Val	Thr	Tyr	Ala
			260					265						270	

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Thr His Arg Ala Ala Glu Thr Val Leu Lys Tyr Tyr Val Lys Gln Leu  
 275 280 285  
 His Ala Leu Glu His Glu Leu Ser Leu Ser Asp Arg Met Asn Val Ile  
 290 295 300  
 Ser Asp Glu Leu Arg Val Leu Ala Asp Ala Gly Gln Asn Asp Met Pro  
 305 310 315 320  
 Ser Arg Val Asp Glu Pro Tyr Arg Arg Ala Ile His Gly Met Arg Gly  
 325 330 335  
 Arg Met Leu Ala Thr Thr Ala Ala Leu Ile Gly Glu Glu Ala Val Glu  
 340 345 350  
 Gly Thr Trp Phe Lys Thr Phe Thr Pro Tyr Thr Asp Thr His Glu Phe  
 355 360 365  
 Lys Arg Asp Leu Asp Ile Val Asp Gly Ser Leu Arg Met Ser Arg Asp  
 370 375 380  
 Asp Ile Ile Ala Asp Asp Arg Leu Ala Met Leu Arg Ser Ala Leu Asp  
 385 390 395 400  
 Ser Phe Gly Phe Asn Leu Tyr Ser Leu Asp Leu Arg Gln Asn Ser Asp  
 405 410 415  
 Gly Phe Glu Asp Val Leu Thr Glu Leu Phe Ala Thr Ala Gln Thr Glu  
 420 425 430  
 Lys Asn Tyr Arg Gly Leu Thr Glu Ala Glu Lys Leu Asp Leu Leu Ile  
 435 440 445  
 Arg Glu Leu Ser Thr Pro Arg Pro Leu Ile Pro His Gly Asp Pro Asp  
 450 455 460  
 Tyr Ser Glu Ala Thr Asn Arg Glu Leu Gly Ile Phe Ser Lys Ala Ala  
 465 470 475 480  
 Glu Ala Val Arg Lys Phe Gly Pro Leu Met Val Pro His Cys Ile Ile  
 485 490 495  
 Ser Met Ala Ser Ser Val Thr Asp Ile Leu Glu Pro Met Val Leu Leu  
 500 505 510  
 Lys Glu Phe Gly Leu Ile Arg Ala Asn Gly Lys Asn Pro Thr Gly Ser  
 515 520 525  
 Val Asp Val Ile Pro Leu Phe Glu Thr Ile Asp Asp Leu Gln Arg Gly  
 530 535 540  
 Ala Gly Ile Leu Glu Glu Leu Trp Asp Ile Asp Leu Tyr Arg Asn Tyr  
 545 550 555 560  
 Leu Glu Gln Arg Asp Asn Val Gln Glu Val Met Leu Gly Tyr Ser Asp  
 565 570 575  
 Ser Asn Lys Asp Gly Gly Tyr Phe Ala Ala Asn Trp Ala Leu Tyr Asp  
 580 585 590  
 Ala Glu Leu Arg Leu Val Glu Leu Cys Arg Gly Arg Asn Val Lys Leu  
 595 600 605  
 Arg Leu Phe His Gly Arg Gly Gly Thr Val Gly Arg Gly Gly Gly Pro

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610	615	620
Ser Tyr Asp Ala Ile Leu	Ala Gln Pro Lys Gly	Ala Val Arg Gly Ala
625	630	635
Val Arg Val Thr Glu	Gln Gly Glu Ile Ile	Ser Ala Lys Tyr Gly Asn
645	650	655
Pro Asp Thr Ala Arg	Arg Asn Leu Glu Ala	Leu Val Ser Ala Thr Leu
660	665	670
Glu Ala Ser Leu Leu	Asp Asp Val Glu Leu	Pro Asn Arg Glu Arg Ala
675	680	685
His Gln Ile Met Gly	Glu Ile Ser Glu Leu	Ser Phe Arg Arg Tyr Ser
690	695	700
Ser Leu Val His Glu	Asp Pro Gly Phe Ile	Gln Tyr Phe Thr Gln Ser
705	710	715
Thr Pro Leu Gln Glu	Ile Gly Ser Leu Asn	Ile Gly Ser Arg Pro Ser
725	730	735
Ser Arg Lys Gln Thr	Asn Thr Val Glu Asp	Leu Arg Ala Ile Pro Trp
740	745	750
Val Leu Ser Trp Ser	Gln Ser Arg Val Met	Leu Pro Gly Trp Phe Gly
755	760	765
Val Gly Thr Ala Leu	Arg Glu Trp Ile Gly	Glu Gly Glu Gly Ala Ala
770	775	780
Glu Arg Ile Ala Glu	Leu Gln Glu Leu Asn	Arg Cys Trp Pro Phe Phe
785	790	795
Thr Ser Val Leu Asp	Asn Met Ala Gln Val	Met Ser Lys Ala Glu Leu
805	810	815
Arg Leu Ala Arg Leu	Tyr Ala Asp Leu Ile	Pro Asp Arg Glu Val Ala
820	825	830
Asp Arg Ile Tyr Glu	Thr Ile Phe Gly Glu	Tyr Phe Leu Thr Lys Glu
835	840	845
Met Phe Cys Thr Ile	Thr Gly Ser Gln Asp	Leu Leu Asp Asp Asn Pro
850	855	860
Ala Leu Ala Arg Ser	Val Arg Ser Arg Phe	Pro Tyr Leu Leu Pro Leu
865	870	875
Asn Val Ile Gln Val	Glu Met Met Arg Arg	Tyr Arg Ser Gly Asp Glu
885	890	895
Gly Thr Ala Val Pro	Arg Asn Ile Arg Leu	Thr Met Asn Gly Leu Ser
900	905	910
Thr Ala Leu Arg Asn	Ser Gly	
915		

&lt;210&gt; 27

&lt;211&gt; 3907

&lt;212&gt; DNA

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<213> *Corynebacterium thermoaminogenes*

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (686)..(3388)

&lt;400&gt; 27

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attacttcag ctgactcagc aacattcgta ttaggtatgc aaacaacatt tggttcgtta 60
aatccaagta gtaatggttaa agtaactlgg ggtatlgctc aagcacitai cgcctllgta 120
tlatiattag ctggtagcgg agatggaact aaagctctca acgcaaitca gagtgccgct 180
attattagtg cgtttccatt cttctllgtc gtcataitaa tgaatgacag tttctacaaa 240
gatgctaata aagaacgiaa attcttagga ttaacattaa cgcctaataa acacagatta 300
gaagaatacg ttaaalaica acaagaggat tacgaatctg atattllaga aaaacglgaa 360
tctagacgia atcgtgaaag agaagaataa ttgaatgaaa tatctactat aatggtaggt 420
ttaaagctat caacaatttt gttagatagct atttttaigt ttcaaacata taaatattat 480
ttactlgcga ttgataacca ttctcaatta ataaaaataa cttataglac aaatgcgtta 540
taataagttt tactataact accgatitaa aaatgcgaaa tgaaaaaatga cccctttata 600
tacctataca gttagtllcg aaaacatata ataatacaat ttaactaagg catataaata 660
talagaaatt caagggggat atcaa atg gct tct aat ttt aaa gaa aca gcg 712
Met Ala Ser Asn Phe Lys Glu Thr Ala
      1              5
aag aaa caa ttt gat tta aat ggc caa tca tac acg tac tat gat tta 760
Lys Lys Gln Phe Asp Leu Asn Gly Gln Ser Tyr Thr Tyr Tyr Asp Leu
      10              15              20              25
aaa tca tta gaa gaa caa ggt tta act aaa att tca aag tta cct tat 808
Lys Ser Leu Glu Glu Gln Gly Leu Thr Lys Ile Ser Lys Leu Pro Tyr
      30              35              40
tca atc cgt gla tta cta gaa tca gtg tta cgt cag gaa gat gat ttt 856
Ser Ile Arg Val Leu Leu Glu Ser Val Leu Arg Gln Glu Asp Asp Phe
      45              50              55
gla att act gat gat cac att aaa caa tta gca gaa ttt ggc aaa aaa 904
Val Ile Thr Asp Asp His Ile Lys Gln Leu Ala Glu Phe Gly Lys Lys
      60              65              70
ggt aac gaa ggt gaa gla cct ttc aaa cca tct cga gtt att tta caa 952
Gly Asn Glu Gly Glu Val Pro Phe Lys Pro Ser Arg Val Ile Leu Gln
      75              80              85
gac ttc act ggt gla cca gca gla gtt gac tta gcg tct tta cgt aaa 1000
Asp Phe Thr Gly Val Pro Ala Val Val Asp Leu Ala Ser Leu Arg Lys
      90              95              100              105
gca atg aat gat gtt ggt ggg gat att aat aaa att aac cct gaa gla 1048
Ala Met Asn Asp Val Gly Gly Asp Ile Asn Lys Ile Asn Pro Glu Val
      110              115              120
cca gtt gac tta gtt att gac cac tct gla caa gla gat agt tat gct 1096

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Pro	Val	Asp	Leu	Val	Ile	Asp	His	Ser	Val	Gln	Val	Asp	Ser	Tyr	Ala	
			125					130				135				
aat	cca	gat	gca	tta	caa	cgt	aac	atg	aaa	tta	gaa	ttt	gaa	cgt	aac	1144
Asn	Pro	Asp	Ala	Leu	Gln	Arg	Asn	Met	Lys	Leu	Glu	Phe	Glu	Arg	Asn	
			140				145					150				
tat	gaa	cgt	tac	caa	ttc	tta	aac	igg	gca	aca	aaa	gca	ttt	gat	aac	1192
Tyr	Glu	Arg	Tyr	Gln	Phe	Leu	Asn	Trp	Ala	Thr	Lys	Ala	Phe	Asp	Asn	
			155				160					165				
tat	aat	gca	gta	cca	ccg	gct	aca	ggg	att	gtc	cac	caa	gta	aac	tta	1240
Tyr	Asn	Ala	Val	Pro	Pro	Ala	Thr	Gly	Ile	Val	His	Gln	Val	Asn	Leu	
			170			175				180					185	
gaa	tac	tta	gcg	aat	gtt	gta	cat	gtt	cgt	gac	gtt	gac	gga	gaa	caa	1288
Glu	Tyr	Leu	Ala	Asn	Val	Val	His	Val	Arg	Asp	Val	Asp	Gly	Glu	Gln	
			190					195				200				
act	gct	ttc	cca	gat	aca	tta	gtt	ggg	act	gac	tca	cat	act	aca	atg	1336
Thr	Ala	Phe	Pro	Asp	Thr	Leu	Val	Gly	Thr	Asp	Ser	His	Thr	Thr	Met	
			205					210				215				
att	aac	ggg	att	ggg	gta	tta	ggg	igg	ggg	gtc	ggc	ggg	atc	gaa	gct	1384
Ile	Asn	Gly	Ile	Gly	Val	Leu	Gly	Trp	Gly	Val	Gly	Gly	Ile	Glu	Ala	
			220				225					230				
gaa	gca	ggg	atg	tta	gga	caa	cca	tca	tac	ttc	cca	att	cca	gaa	gtt	1432
Glu	Ala	Gly	Met	Leu	Gly	Gln	Pro	Ser	Tyr	Phe	Pro	Ile	Pro	Glu	Val	
			235			240					245					
att	ggg	gtt	aaa	tta	agt	aat	gaa	tta	cca	caa	ggg	tca	aca	gca	act	1480
Ile	Gly	Val	Lys	Leu	Ser	Asn	Glu	Leu	Pro	Gln	Gly	Ser	Thr	Ala	Thr	
			250			255				260					265	
gac	tta	gca	tta	cgt	gta	act	gaa	gag	tta	cgt	aaa	cgt	ggg	gta	gta	1528
Asp	Leu	Ala	Leu	Arg	Val	Thr	Glu	Glu	Leu	Arg	Lys	Arg	Gly	Val	Val	
			270					275				280				
ggg	aaa	ttc	gtt	gag	ttc	ttt	ggg	ccg	ggg	gta	aca	aac	tta	cca	tta	1576
Gly	Lys	Phe	Val	Glu	Phe	Phe	Gly	Pro	Gly	Val	Thr	Asn	Leu	Pro	Leu	
			285				290					295				
gct	gac	cgt	gca	aca	att	gcg	aac	atg	gcg	ccg	gaa	tat	ggg	gca	act	1624
Ala	Asp	Arg	Ala	Thr	Ile	Ala	Asn	Met	Ala	Pro	Glu	Tyr	Gly	Ala	Thr	
			300				305					310				
tgt	ggg	ttc	ttc	cca	gtt	gat	gaa	gaa	tca	ctt	aaa	tac	atg	aaa	tta	1672
Cys	Gly	Phe	Phe	Pro	Val	Asp	Glu	Glu	Ser	Leu	Lys	Tyr	Met	Lys	Leu	
			315			320					325					
act	ggg	cgt	aaa	gat	gat	cat	att	gca	cta	gta	aaa	gaa	tat	tta	caa	1720
Thr	Gly	Arg	Lys	Asp	Asp	His	Ile	Ala	Leu	Val	Lys	Glu	Tyr	Leu	Gln	
			330			335				340					345	
caa	aat	aat	atg	ttc	ttc	caa	gtt	gaa	aat	gaa	gat	ccg	gaa	tat	act	1768
Gln	Asn	Asn	Met	Phe	Phe	Gln	Val	Glu	Asn	Glu	Asp	Pro	Glu	Tyr	Thr	



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															350																355																360	
gaa	gtg	att	gat	tta	gat	tta	tct	aca	ggt	caa	gct	tct	tta	tca	ggt	1816	Glu	Val	Ile	Asp	Leu	Asp	Leu	Ser	Thr	Val	Gln	Ala	Ser	Leu	Ser	Gly																
															365																370																375	
cca	aaa	cgt	cca	caa	gat	tta	atc	ttc	tta	agt	gac	atg	aaa	act	gaa	1864	Pro	Lys	Arg	Pro	Gln	Asp	Leu	Ile	Phe	Leu	Ser	Asp	Met	Lys	Thr	Glu																
															380																385																390	
ttc	gaa	aaa	tca	gtt	aca	gca	cca	gct	ggt	aac	caa	ggt	cac	ggt	tta	1912	Phe	Glu	Lys	Ser	Val	Thr	Ala	Pro	Ala	Gly	Asn	Gln	Gly	His	Gly	Leu																
															395																400																405	
gat	gaa	agt	gaa	ttt	gat	aag	aaa	gca	gaa	atc	aaa	ttt	aat	gat	ggt	1960	Asp	Glu	Ser	Glu	Phe	Asp	Lys	Lys	Ala	Glu	Ile	Lys	Phe	Asn	Asp	Gly																
															410																415																420	425
aga	act	tca	act	atg	aag	act	ggt	gat	gtt	gcg	att	gca	gcg	att	aca	2008	Arg	Thr	Ser	Thr	Met	Lys	Thr	Gly	Asp	Val	Ala	Ile	Ala	Ala	Ile	Thr																
															430																435																440	
tca	tgt	aca	aat	aca	tct	aac	cct	tac	gtt	atg	tta	ggt	gca	ggt	tta	2056	Ser	Cys	Thr	Asn	Thr	Ser	Asn	Pro	Tyr	Val	Met	Leu	Gly	Ala	Gly	Leu																
															445																450																455	
gta	gct	aaa	aaa	gca	att	gaa	aaa	ggc	tta	aaa	gta	cct	gat	tat	gta	2104	Val	Ala	Lys	Lys	Ala	Ile	Glu	Lys	Gly	Leu	Lys	Val	Pro	Asp	Tyr	Val																
															460																465																470	
aaa	act	tca	tta	gca	cca	ggt	tca	aaa	gtt	gtt	act	gga	tat	tta	aga	2152	Lys	Thr	Ser	Leu	Ala	Pro	Gly	Ser	Lys	Val	Val	Thr	Gly	Tyr	Leu	Arg																
															475																480																485	
gat	tca	ggt	tta	caa	gaa	tat	cct	gat	gac	tta	ggt	ttc	aac	tta	gtt	2200	Asp	Ser	Gly	Leu	Gln	Glu	Tyr	Leu	Asp	Asp	Leu	Gly	Phe	Asn	Leu	Val																
															490																495																500	505
ggt	tat	ggt	tgt	aca	act	tgt	atc	ggt	aac	tca	ggt	cca	tta	tta	cct	2248	Gly	Tyr	Gly	Cys	Thr	Thr	Cys	Ile	Gly	Asn	Ser	Gly	Pro	Leu	Leu	Pro																
															510																515																520	
gaa	att	gaa	aaa	gca	gta	gct	gac	gaa	gat	tta	tta	gta	act	tct	gta	2296	Glu	Ile	Glu	Lys	Ala	Val	Ala	Asp	Glu	Asp	Leu	Leu	Val	Thr	Ser	Val																
															525																530																535	
cct	tct	ggt	aac	cgt	aac	ttt	gaa	ggt	cgt	atc	cat	ccg	tta	gtt	aaa	2344	Leu	Ser	Gly	Asn	Arg	Asn	Phe	Glu	Gly	Arg	Ile	His	Pro	Leu	Val	Lys																
															540																545																550	
gct	aac	tac	tta	gct	tca	cca	caa	tta	gtt	gta	gct	tat	gca	tta	gct	2392	Ala	Asn	Tyr	Leu	Ala	Ser	Pro	Gln	Leu	Val	Val	Ala	Tyr	Ala	Leu	Ala																
															555																560																565	
gga	acg	gtt	gat	atc	gat	tta	cac	aat	gaa	cct	atc	ggt	aaa	ggt	aaa	2440	Gly	Thr	Val	Asp	Ile	Asp	Leu	His	Asn	Glu	Pro	Ile	Gly	Lys	Gly	Lys																
															570																575																580	585

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gat ggc gaa gat gta tac ctt aaa gat atc tgg cca agt atc aaa gaa	2488
Asp Gly Glu Asp Val Tyr Leu Lys Asp Ile Trp Pro Ser Ile Lys Glu	
590 595 600	
gtt gca gac act gtt gat agt gtc gta acg cca gaa tta ttc tta gaa	2536
Val Ala Asp Thr Val Asp Ser Val Val Thr Pro Glu Leu Phe Leu Glu	
605 610 615	
gaa tat gca aat gta tac gaa aat aat gaa atg tgg aat gaa atc gac	2584
Glu Tyr Ala Asn Val Tyr Glu Asn Asn Glu Met Trp Asn Glu Ile Asp	
620 625 630	
gtt act gac gca cca tta tat gat ttc gat cca aat tca act tat att	2632
Val Thr Asp Ala Pro Leu Tyr Asp Phe Asp Pro Asn Ser Thr Tyr Ile	
635 640 645	
caa aat cca tca ttc ttc caa ggt tta tct aaa gaa cca gga act att	2680
Gln Asn Pro Ser Phe Phe Gln Gly Leu Ser Lys Glu Pro Gly Thr Ile	
650 655 660 665	
gaa cca tta aaa gat tta cgt att atg ggt aaa ttt ggt gat tca gtt	2728
Glu Pro Leu Lys Asp Leu Arg Ile Met Gly Lys Phe Gly Asp Ser Val	
670 675 680	
aca act gac cac att tct cca gca ggt gcg atc ggt aaa gat aca cca	2776
Thr Thr Asp His Ile Ser Pro Ala Gly Ala Ile Gly Lys Asp Thr Pro	
685 690 695	
gca ggt aaa tat tta tta gac cat gat gtt cca att aga gaa ttt aac	2824
Ala Gly Lys Tyr Leu Leu Asp His Asp Val Pro Ile Arg Glu Phe Asn	
700 705 710	
tct tat ggt tca aga cgt ggt aac cat gaa gta atg gta cgt ggt act	2872
Ser Tyr Gly Ser Arg Arg Gly Asn His Glu Val Met Val Arg Gly Thr	
715 720 725	
ttc gct aat atc cgt att aaa aac caa tta gca cca ggc act gaa ggt	2920
Phe Ala Asn Ile Arg Ile Lys Asn Gln Leu Ala Pro Gly Thr Glu Gly	
730 735 740 745	
gga ttt aca aca tat tgg cct aca gaa gaa atc atg cct atc tat gat	2968
Gly Phe Thr Thr Tyr Trp Pro Thr Glu Glu Ile Met Pro Ile Tyr Asp	
750 755 760	
gca gct atg aga tac aaa gaa aat ggt act ggt tta gct gtt tta gct	3016
Ala Ala Met Arg Tyr Lys Glu Asn Gly Thr Gly Leu Ala Val Leu Ala	
765 770 775	
ggt aat gat tac ggt atg ggt tca tct cgt gac tgg gca gct aaa ggt	3064
Gly Asn Asp Tyr Gly Met Gly Ser Ser Arg Asp Trp Ala Ala Lys Gly	
780 785 790	
act aac tta tta ggt gtt aaa act gtt att gca caa agt tat gaa cgt	3112
Thr Asn Leu Leu Gly Val Lys Thr Val Ile Ala Gln Ser Tyr Glu Arg	
795 800 805	
atc cat cgt tca aac tta gta atg atg ggt gta tta cca tta caa ttt	3160

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Ile His Arg Ser Asn Leu Val Met Met Gly Val Leu Pro Leu Gln Phe  
 810 815 820 825  
 aaa caa ggt gag tca gct gat tct cta ggt tta gaa ggt aaa gaa gaa 3208  
 Lys Gln Gly Glu Ser Ala Asp Ser Leu Gly Leu Glu Gly Lys Glu Glu  
 830 835 840  
 att tct gta gat atc gat gaa aat gtt aaa cca cat gat tta gta act 3256  
 Ile Ser Val Asp Ile Asp Glu Asn Val Lys Pro His Asp Leu Val Thr  
 845 850 855  
 gtt cat gct aaa aaa gaa aac gga gaa gtt gtt gat ttt gaa gca atg 3304  
 Val His Ala Lys Lys Glu Asn Gly Glu Val Val Asp Phe Glu Ala Met  
 860 865 870  
 gtt cgt ttc gat tca tta gta gaa tta gat tat tat cgt cat ggt ggt 3352  
 Val Arg Phe Asp Ser Leu Val Glu Leu Asp Tyr Tyr Arg His Gly Gly  
 875 880 885  
 atc tta caa atg gta tta aga aac aaa tta gct caa taatcacaat 3398  
 Ile Leu Gln Met Val Leu Arg Asn Lys Leu Ala Gln  
 890 895 900  
 gtgacititg acagtgctaa cgtiltaggtt agcacigtitl tittatgctt aaclatata 3458  
 gtaatgttaa tagttaagga aggatiggac tlaaatgatt tatagtttga ctgaaattga 3518  
 accaagatai caagagacag ataaaatggg cgtgatitai catggcaatt atgcaacaig 3578  
 gtttgaagta gcgcgtacag attacattag aaaactagga tttagttatg ctgatatgga 3638  
 aaagcaaggg aicatttctc cagttacaga cttaaataic aaatataaaa aatcaatttt 3698  
 ttatcctgaa aaagtaacca ttaaaacaig ggttgaaaaa taitcaagal tacgttctgt 3758  
 glatagatai gaaatttita atgaacaggg agaacttgca actacaggtt atactgagtt 3818  
 aatttgiatg aaagctgata cctttagacc aattagattt gatcgttatt tctcagattg 3878  
 gcatgaaacc tatagtaaag ttgaagctt 3907

&lt;210&gt; 28

&lt;211&gt; 901

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 28

Met Ala Ser Asn Phe Lys Glu Thr Ala Lys Lys Gln Phe Asp Leu Asn  
 1 5 10 15  
 Gly Gln Ser Tyr Thr Tyr Tyr Asp Leu Lys Ser Leu Glu Glu Gln Gly  
 20 25 30  
 Leu Thr Lys Ile Ser Lys Leu Pro Tyr Ser Ile Arg Val Leu Leu Glu  
 35 40 45  
 Ser Val Leu Arg Gln Glu Asp Asp Phe Val Ile Thr Asp Asp His Ile  
 50 55 60  
 Lys Gln Leu Ala Glu Phe Gly Lys Lys Gly Asn Glu Gly Glu Val Pro  
 65 70 75 80

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Phe	Lys	Pro	Ser	Arg	Val	Ile	Leu	Gln	Asp	Phe	Thr	Gly	Val	Pro	Ala	
				85					90					95		
Val	Val	Asp	Leu	Ala	Ser	Leu	Arg	Lys	Ala	Met	Asn	Asp	Val	Gly	Gly	
			100					105					110			
Asp	Ile	Asn	Lys	Ile	Asn	Pro	Glu	Val	Pro	Val	Asp	Leu	Val	Ile	Asp	
		115					120					125				
His	Ser	Val	Gln	Val	Asp	Ser	Tyr	Ala	Asn	Pro	Asp	Ala	Leu	Gln	Arg	
	130					135					140					
Asn	Met	Lys	Leu	Glu	Phe	Glu	Arg	Asn	Tyr	Glu	Arg	Tyr	Gln	Phe	Leu	
145					150					155					160	
Asn	Trp	Ala	Thr	Lys	Ala	Phe	Asp	Asn	Tyr	Asn	Ala	Val	Pro	Pro	Ala	
				165					170						175	
Thr	Gly	Ile	Val	His	Gln	Val	Asn	Leu	Glu	Tyr	Leu	Ala	Asn	Val	Val	
			180					185						190		
His	Val	Arg	Asp	Val	Asp	Gly	Glu	Gln	Thr	Ala	Phe	Pro	Asp	Thr	Leu	
		195					200					205				
Val	Gly	Thr	Asp	Ser	His	Thr	Thr	Met	Ile	Asn	Gly	Ile	Gly	Val	Leu	
	210					215					220					
Gly	Trp	Gly	Val	Gly	Gly	Ile	Glu	Ala	Glu	Ala	Gly	Met	Leu	Gly	Gln	
225					230					235					240	
Pro	Ser	Tyr	Phe	Pro	Ile	Pro	Glu	Val	Ile	Gly	Val	Lys	Leu	Ser	Asn	
				245					250					255		
Glu	Leu	Pro	Gln	Gly	Ser	Thr	Ala	Thr	Asp	Leu	Ala	Leu	Arg	Val	Thr	
			260					265					270			
Glu	Glu	Leu	Arg	Lys	Arg	Gly	Val	Val	Gly	Lys	Phe	Val	Glu	Phe	Phe	
		275					280					285				
Gly	Pro	Gly	Val	Thr	Asn	Leu	Pro	Leu	Ala	Asp	Arg	Ala	Thr	Ile	Ala	
	290					295					300					
Asn	Met	Ala	Pro	Glu	Tyr	Gly	Ala	Thr	Cys	Gly	Phe	Phe	Pro	Val	Asp	
305					310					315					320	
Glu	Glu	Ser	Leu	Lys	Tyr	Met	Lys	Leu	Thr	Gly	Arg	Lys	Asp	Asp	His	
				325					330					335		
Ile	Ala	Leu	Val	Lys	Glu	Tyr	Leu	Gln	Gln	Asn	Asn	Met	Phe	Phe	Gln	
			340					345					350			
Val	Glu	Asn	Glu	Asp	Pro	Glu	Tyr	Thr	Glu	Val	Ile	Asp	Leu	Asp	Leu	
		355					360					365				
Ser	Thr	Val	Gln	Ala	Ser	Leu	Ser	Gly	Pro	Lys	Arg	Pro	Gln	Asp	Leu	
	370					375					380					
Ile	Phe	Leu	Ser	Asp	Met	Lys	Thr	Glu	Phe	Glu	Lys	Ser	Val	Thr	Ala	
385					390					395					400	
Pro	Ala	Gly	Asn	Gln	Gly	His	Gly	Leu	Asp	Glu	Ser	Glu	Phe	Asp	Lys	
				405					410					415		
Lys	Ala	Glu	Ile	Lys	Phe	Asn	Asp	Gly	Arg	Thr	Ser	Thr	Met	Lys	Thr	

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			420					425					430			
Gly	Asp	Val	Ala	Ile	Ala	Ala	Ile	Thr	Ser	Cys	Thr	Asn	Thr	Ser	Asn	
		435					440					445				
Pro	Tyr	Val	Met	Leu	Gly	Ala	Gly	Leu	Val	Ala	Lys	Lys	Ala	Ile	Glu	
	450					455					460					
Lys	Gly	Leu	Lys	Val	Pro	Asp	Tyr	Val	Lys	Thr	Ser	Leu	Ala	Pro	Gly	
465					470					475					480	
Ser	Lys	Val	Val	Thr	Gly	Tyr	Leu	Arg	Asp	Ser	Gly	Leu	Gln	Glu	Tyr	
			485					490						495		
Leu	Asp	Asp	Leu	Gly	Phe	Asn	Leu	Val	Gly	Tyr	Gly	Cys	Thr	Thr	Cys	
		500					505						510			
Ile	Gly	Asn	Ser	Gly	Pro	Leu	Leu	Pro	Glu	Ile	Glu	Lys	Ala	Val	Ala	
	515					520					525					
Asp	Glu	Asp	Leu	Leu	Val	Thr	Ser	Val	Leu	Ser	Gly	Asn	Arg	Asn	Phe	
	530					535				540						
Glu	Gly	Arg	Ile	His	Pro	Leu	Val	Lys	Ala	Asn	Tyr	Leu	Ala	Ser	Pro	
545					550					555					560	
Gln	Leu	Val	Val	Ala	Tyr	Ala	Leu	Ala	Gly	Thr	Val	Asp	Ile	Asp	Leu	
			565					570						575		
His	Asn	Glu	Pro	Ile	Gly	Lys	Gly	Lys	Asp	Gly	Glu	Asp	Val	Tyr	Leu	
		580					585						590			
Lys	Asp	Ile	Trp	Pro	Ser	Ile	Lys	Glu	Val	Ala	Asp	Thr	Val	Asp	Ser	
	595					600					605					
Val	Val	Thr	Pro	Glu	Leu	Phe	Leu	Glu	Glu	Tyr	Ala	Asn	Val	Tyr	Glu	
	610					615					620					
Asn	Asn	Glu	Met	Trp	Asn	Glu	Ile	Asp	Val	Thr	Asp	Ala	Pro	Leu	Tyr	
625					630					635					640	
Asp	Phe	Asp	Pro	Asn	Ser	Thr	Tyr	Ile	Gln	Asn	Pro	Ser	Phe	Phe	Gln	
			645					650						655		
Gly	Leu	Ser	Lys	Glu	Pro	Gly	Thr	Ile	Glu	Pro	Leu	Lys	Asp	Leu	Arg	
		660					665						670			
Ile	Met	Gly	Lys	Phe	Gly	Asp	Ser	Val	Thr	Thr	Asp	His	Ile	Ser	Pro	
	675					680						685				
Ala	Gly	Ala	Ile	Gly	Lys	Asp	Thr	Pro	Ala	Gly	Lys	Tyr	Leu	Leu	Asp	
	690					695					700					
His	Asp	Val	Pro	Ile	Arg	Glu	Phe	Asn	Ser	Tyr	Gly	Ser	Arg	Arg	Gly	
705					710					715					720	
Asn	His	Glu	Val	Met	Val	Arg	Gly	Thr	Phe	Ala	Asn	Ile	Arg	Ile	Lys	
			725					730						735		
Asn	Gln	Leu	Ala	Pro	Gly	Thr	Glu	Gly	Gly	Phe	Thr	Thr	Tyr	Trp	Pro	
		740					745									

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Asn Gly Thr Gly Leu Ala Val Leu Ala Gly Asn Asp Tyr Gly Met Gly  
 770 775 780  
 Ser Ser Arg Asp Trp Ala Ala Lys Gly Thr Asn Leu Leu Gly Val Lys  
 785 790 795 800  
 Thr Val Ile Ala Gln Ser Tyr Glu Arg Ile His Arg Ser Asn Leu Val  
 805 810 815  
 Met Met Gly Val Leu Pro Leu Gln Phe Lys Gln Gly Glu Ser Ala Asp  
 820 825 830  
 Ser Leu Gly Leu Glu Gly Lys Glu Glu Ile Ser Val Asp Ile Asp Glu  
 835 840 845  
 Asn Val Lys Pro His Asp Leu Val Thr Val His Ala Lys Lys Glu Asn  
 850 855 860  
 Gly Glu Val Val Asp Phe Glu Ala Met Val Arg Phe Asp Ser Leu Val  
 865 870 875 880  
 Glu Leu Asp Tyr Tyr Arg His Gly Gly Ile Leu Gln Met Val Leu Arg  
 885 890 895  
 Asn Lys Leu Ala Gln  
 900

&lt;210&gt; 29

&lt;211&gt; 3006

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (328)..(2514)

&lt;400&gt; 29

gtcgacgacg aacccccac cgccgaacca gccgccgac tgggtgtggga gacacccggg 60  
 ttctctctcc tgggtgaaca ggigccacaa ccccgiccca acaggcacac ctaccactgg 120  
 atcgccgggg agagcagcat ggtcacacgc ctgcggcggt ccttggtgaa ggatcacggc 180  
 ctggacagat cgcaggtggc attcatgggt tatlggaggc agggagtggc catgaggggt 240  
 tgatacgct tccctgaggg tccgcaggcg tgcctcacc tgtattcttg atagtgaac 300  
 aaaagagccc acataacaag gagactc atg gct aag atc atc tgg acc cgc acc 354  
 Met Ala Lys Ile Ile Trp Thr Arg Thr  
 1 5  
 gac gaa gca ccg ctg ctc gcg acc tac tcg ctg aag ccg gtc gtc gag 402  
 Asp Glu Ala Pro Leu Leu Ala Thr Tyr Ser Leu Lys Pro Val Val Glu  
 10 15 20 25  
 gct ttc gcc gcc acc gcg ggc atc gag gtg gag acc cgc gat atc tct 450  
 Ala Phe Ala Ala Thr Ala Gly Ile Glu Val Glu Thr Arg Asp Ile Ser  
 30 35 40

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ctc gcc ggt cgc atc ctc gca cag ttc gcg gac cag ctc ccc gag gag	498
Leu Ala Gly Arg Ile Leu Ala Gln Phe Ala Asp Gln Leu Pro Glu Glu	
45 50 55	
cag aag gtc tcc gac gcc ctc gcc gag ctc ggc gaa ctg gct aag acc	546
Gln Lys Val Ser Asp Ala Leu Ala Glu Leu Gly Glu Leu Ala Lys Thr	
60 65 70	
ccc gaa gcc aac atc atc aag ctt ccc aac atc tcc gca tcc gta ccg	594
Pro Glu Ala Asn Ile Ile Lys Leu Pro Asn Ile Ser Ala Ser Val Pro	
75 80 85	
cag ctc aag gct gcc gta aag gaa ctg cag gaa cag ggc tac gac ctg	642
Gln Leu Lys Ala Ala Val Lys Glu Leu Gln Glu Gln Gly Tyr Asp Leu	
90 95 100 105	
ccc gag tac gag gat gcc aag gac cgc tac gcc gct gtc atc ggc tcc	690
Pro Glu Tyr Glu Asp Ala Lys Asp Arg Tyr Ala Ala Val Ile Gly Ser	
110 115 120	
aac gtc aac ccg gtc ctg cgc gag ggc aac tcc gac cgc cgc gca ccg	738
Asn Val Asn Pro Val Leu Arg Glu Gly Asn Ser Asp Arg Arg Ala Pro	
125 130 135	
gtg gcc gtg aag aac ttc gtg aag aag ttc ccc cac cgc atg ggc gag	786
Val Ala Val Lys Asn Phe Val Lys Lys Phe Pro His Arg Met Gly Glu	
140 145 150	
tgg tcc gcc gac tcc aag acc aac gtt gcc acc atg ggt gcc gac gac	834
Trp Ser Ala Asp Ser Lys Thr Asn Val Ala Thr Met Gly Ala Asp Asp	
155 160 165	
ttc cgc agc aat gag aag tcc gtg atc atg gac gag gcc gac acc gtg	882
Phe Arg Ser Asn Glu Lys Ser Val Ile Met Asp Glu Ala Asp Thr Val	
170 175 180 185	
gtg atc aag cat gtc gcc gcc gac ggc acc gag acc gtg ctc aag gac	930
Val Ile Lys His Val Ala Ala Asp Gly Thr Glu Thr Val Leu Lys Asp	
190 195 200	
agc ctc ccc ctg ctc aag ggt gag gtc atc gac ggc acc ttc atc tcc	978
Ser Leu Pro Leu Leu Lys Gly Glu Val Ile Asp Gly Thr Phe Ile Ser	
205 210 215	
gcc aag gca ctg gac gcc ttc ctg ctc gac cag gtc aaa cgc gcc aag	1026
Ala Lys Ala Leu Asp Ala Phe Leu Leu Asp Gln Val Lys Arg Ala Lys	
220 225 230	
gag gag ggc atc ctc ttc tcc gcc cac atg aag gcc acc atg atg aag	1074
Glu Glu Gly Ile Leu Phe Ser Ala His Met Lys Ala Thr Met Met Lys	
235 240 245	
gtc tcc gac ccg atc atc ttc ggc cac atc gtc cgc gcc tac ttc gcc	1122
Val Ser Asp Pro Ile Ile Phe Gly His Ile Val Arg Ala Tyr Phe Ala	
250 255 260 265	
gat gtc tac gca cag tac ggt gag cag ctg ctc gcc gcc ggc ctc aac	1170

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Asp	Val	Tyr	Ala	Gln	Tyr	Gly	Glu	Gln	Leu	Leu	Ala	Ala	Gly	Leu	Asn	
				270					275					280		
ggt	gag	aac	ggt	ctc	gcc	gcc	atc	tac	gcc	ggc	ctg	gac	aag	ctg	gac	1218
Gly	Glu	Asn	Gly	Leu	Ala	Ala	Ile	Tyr	Ala	Gly	Leu	Asp	Lys	Leu	Asp	
			285					290					295			
aac	ggt	gcc	gag	atc	aag	gca	gcc	ttc	gac	aag	ggc	ctg	gaa	gag	ggc	1266
Asn	Gly	Ala	Glu	Ile	Lys	Ala	Ala	Phe	Asp	Lys	Gly	Leu	Glu	Glu	Gly	
		300					305					310				
ccc	gac	ctg	gcc	atg	gtg	aac	tcc	gcc	aag	ggc	atc	acc	aac	ctg	cat	1314
Pro	Asp	Leu	Ala	Met	Val	Asn	Ser	Ala	Lys	Gly	Ile	Thr	Asn	Leu	His	
		315				320					325					
gtg	ccc	tcc	gat	gtc	atc	atc	gac	gcc	tcc	atg	ccc	gcc	atg	atc	cgc	1362
Val	Pro	Ser	Asp	Val	Ile	Ile	Asp	Ala	Ser	Met	Pro	Ala	Met	Ile	Arg	
330					335				340					345		
acc	tcc	ggc	aag	atg	tgg	aac	aag	gac	gac	cag	acc	cag	gat	gcc	ctg	1410
Thr	Ser	Gly	Lys	Met	Trp	Asn	Lys	Asp	Asp	Gln	Thr	Gln	Asp	Ala	Leu	
			350					355					360			
gct	gtc	atc	ccg	gac	tcc	tcc	tac	gcc	ggt	gtc	tac	cag	acc	gtc	atc	1458
Ala	Val	Ile	Pro	Asp	Ser	Ser	Tyr	Ala	Gly	Val	Tyr	Gln	Thr	Val	Ile	
		365						370				375				
gag	gac	tgc	cgc	aag	aat	ggc	gcc	ttc	gat	ccg	acc	acc	atg	ggc	acc	1506
Glu	Asp	Cys	Arg	Lys	Asn	Gly	Ala	Phe	Asp	Pro	Thr	Thr	Met	Gly	Thr	
		380				385					390					
gtc	ccc	aac	gtc	ggt	ctg	atg	gca	cag	aag	gcc	gag	gag	tac	ggc	tcc	1554
Val	Pro	Asn	Val	Gly	Leu	Met	Ala	Gln	Lys	Ala	Glu	Glu	Tyr	Gly	Ser	
		395			400						405					
cac	gac	aag	acc	ttc	cgt	atc	gag	gcc	gac	ggc	aag	gta	cag	gtc	gtc	1602
His	Asp	Lys	Thr	Phe	Arg	Ile	Glu	Ala	Asp	Gly	Lys	Val	Gln	Val	Val	
410					415				420					425		
gcc	tcc	aac	ggt	gat	gtc	ctc	atc	gag	cac	gac	gtg	gag	aag	ggc	gac	1650
Ala	Ser	Asn	Gly	Asp	Val	Leu	Ile	Glu	His	Asp	Val	Glu	Lys	Gly	Asp	
			430					435				440				
atc	tgg	cgc	gcc	tgc	cag	acc	aag	gac	gcc	ccg	atc	cag	gac	tgg	gtc	1698
Ile	Trp	Arg	Ala	Cys	Gln	Thr	Lys	Asp	Ala	Pro	Ile	Gln	Asp	Trp	Val	
		445						450				455				
aag	ctg	gct	gtc	aac	cgc	gca	cgt	ctc	tcc	ggc	atg	ccc	gct	gtg	ttc	1746
Lys	Leu	Ala	Val	Asn	Arg	Ala	Arg	Leu	Ser	Gly	Met	Pro	Ala	Val	Phe	
		460					465				470					
tgg	ctg	gat	ccc	gcc	cgc	gca	cac	gac	cgc	aac	ctg	acc	aca	ctg	gtg	1794
Trp	Leu	Asp	Pro	Ala	Arg	Ala	His	Asp	Arg	Asn	Leu	Thr	Thr	Leu	Val	
		475				480					485					
gag	aag	tac	ctg	gca	gac	cac	gac	acc	gag	ggc	ctg	gac	atc	cag	atc	1842
Glu	Lys	Tyr	Leu	Ala	Asp	His	Asp	Thr	Glu	Gly	Leu	Asp	Ile	Gln	Ile	



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490		495		500		505	
ctc tcc ccc gtc gag gcc acc cag cac gcc atc gac cgc atc cgc cgc	1890						
Leu Ser Pro Val Glu Ala Thr Gln His Ala Ile Asp Arg Ile Arg Arg							
		510		515		520	
ggc gag gac acc atc tcc gtc acc ggt aac gtc ctg cgt gac tac aac	1938						
Gly Glu Asp Thr Ile Ser Val Thr Gly Asn Val Leu Arg Asp Tyr Asn							
		525		530		535	
acc gac ctc ttc ccg atc ctc gag ctg ggc acc tcc gcc aag atg ctc	1986						
Thr Asp Leu Phe Pro Ile Leu Glu Leu Gly Thr Ser Ala Lys Met Leu							
		540		545		550	
tcc gtc gtg cca ctg atg gcc ggc ggt gga ctc ttc gag acc ggt gcc	2034						
Ser Val Val Pro Leu Met Ala Gly Gly Gly Leu Phe Glu Thr Gly Ala							
		555		560		565	
ggt ggc tcc gcc ccg aag cac gtc cag cag gtc atc gag gaa aac cac	2082						
Gly Gly Ser Ala Pro Lys His Val Gln Gln Val Ile Glu Glu Asn His							
		570		575		580	
ctg cgc tgg gat tcc ctc ggt gag ttc ctg gcc ctg gcc gag tcc ttc	2130						
Leu Arg Trp Asp Ser Leu Gly Glu Phe Leu Ala Leu Ala Glu Ser Phe							
		590		595		600	
cgc cac gag ctc aac acc cgc aac aac acc aag gcc ggt gtc ctc gcc	2178						
Arg His Glu Leu Asn Thr Arg Asn Asn Thr Lys Ala Gly Val Leu Ala							
		605		610		615	
gat gcc ctg gac cgt gcg acc gag aag ctc ctc aac gag gag aag tcc	2226						
Asp Ala Leu Asp Arg Ala Thr Glu Lys Leu Leu Asn Glu Glu Lys Ser							
		620		625		630	
ccg tcc cgc aag gtc ggc gag atc gac aac cgt ggt tcc cac ttc tgg	2274						
Pro Ser Arg Lys Val Gly Glu Ile Asp Asn Arg Gly Ser His Phe Trp							
		635		640		645	
ctg gcc acc tac tgg gcc gat gaa ctg gcc aac cag acc gag gac gcc	2322						
Leu Ala Thr Tyr Trp Ala Asp Glu Leu Ala Asn Gln Thr Glu Asp Ala							
		650		655		660	
gag ctg gct gag acc ttc gcc cct gtc gcc gag gcc ctg aac aac cag	2370						
Glu Leu Ala Glu Thr Phe Ala Pro Val Ala Glu Ala Leu Asn Asn Gln							
		670		675		680	
gct gcc gac atc gac gca gca ctc atc ggt gag cag ggc aag cct gtc	2418						
Ala Ala Asp Ile Asp Ala Ala Leu Ile Gly Glu Gln Gly Lys Pro Val							
		685		690		695	
gac ctg ggt ggc tac tac gca ccc tcc gat gag aag acc tcc gcg atc	2466						
Asp Leu Gly Gly Tyr Tyr Ala Pro Ser Asp Glu Lys Thr Ser Ala Ile							
		700		705		710	
atg cgc ccg gtg gcc gca ttc aac gag atc atc gac tcc ctg aag aag	2514						
Met Arg Pro Val Ala Ala Phe Asn Glu Ile Ile Asp Ser Leu Lys Lys							
		715		720		725	

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taaccccttc tccggagccg acagccgacg gccacgctcc cccgcccacg ggggatcgtg 2574
gccgtcggcc gtttctggca ctggagtga cacttcggtg ataatggiga gatgaacagc 2634
ccccgtgtcc ccgccatcct gtccgccgtt tccgccgtgg gtctgatcgc tgcgtgggc 2694
acccccgttg ccgtcgcaga caccatcacc gcggacaccg accgggaaac ctgcgtggcc 2754
agccagaatg acaactccag cgtgatcagg ttctgggaig acctggaggc cgaatgtccgt 2814
gagcagcgcc tgaccgaact ggatgcacag gaccccggcc tcaagaacga catcgaggcc 2874
ttcatcgccg aggaccgggt agccccctcc gcagccgatc tccagagacg gctggatgca 2934
aatgacgccg gtgagggcct ggccaigtg ctacctgaat cccgcaccga ccccgagggtg 2994
gtggacctgc ag 3006

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&lt;210&gt; 30

&lt;211&gt; 729

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 30

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Met Ala Lys Ile Ile Trp Thr Arg Thr Asp Glu Ala Pro Leu Leu Ala
  1              5              10              15
Thr Tyr Ser Leu Lys Pro Val Val Glu Ala Phe Ala Ala Thr Ala Gly
              20              25              30
Ile Glu Val Glu Thr Arg Asp Ile Ser Leu Ala Gly Arg Ile Leu Ala
              35              40              45
Gln Phe Ala Asp Gln Leu Pro Glu Glu Gln Lys Val Ser Asp Ala Leu
              50              55              60
Ala Glu Leu Gly Glu Leu Ala Lys Thr Pro Glu Ala Asn Ile Ile Lys
              65              70              75              80
Leu Pro Asn Ile Ser Ala Ser Val Pro Gln Leu Lys Ala Ala Val Lys
              85              90              95
Glu Leu Gln Glu Gln Gly Tyr Asp Leu Pro Glu Tyr Glu Asp Ala Lys
              100             105             110
Asp Arg Tyr Ala Ala Val Ile Gly Ser Asn Val Asn Pro Val Leu Arg
              115             120             125
Glu Gly Asn Ser Asp Arg Arg Ala Pro Val Ala Val Lys Asn Phe Val
              130             135             140
Lys Lys Phe Pro His Arg Met Gly Glu Trp Ser Ala Asp Ser Lys Thr
              145             150             155             160
Asn Val Ala Thr Met Gly Ala Asp Asp Phe Arg Ser Asn Glu Lys Ser
              165             170             175
Val Ile Met Asp Glu Ala Asp Thr Val Val Ile Lys His Val Ala Ala
              180             185             190
Asp Gly Thr Glu Thr Val Leu Lys Asp Ser Leu Pro Leu Leu Lys Gly
              195             200             205
Glu Val Ile Asp Gly Thr Phe Ile Ser Ala Lys Ala Leu Asp Ala Phe

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210	215	220
Leu Leu Asp Gln Val Lys Arg Ala Lys Glu Glu Gly Ile Leu Phe Ser		
225	230	235
Ala His Met Lys Ala Thr Met Met Lys Val Ser Asp Pro Ile Ile Phe		240
	245	250
Gly His Ile Val Arg Ala Tyr Phe Ala Asp Val Tyr Ala Gln Tyr Gly		255
	260	265
Glu Gln Leu Leu Ala Ala Gly Leu Asn Gly Glu Asn Gly Leu Ala Ala		270
	275	280
Ile Tyr Ala Gly Leu Asp Lys Leu Asp Asn Gly Ala Glu Ile Lys Ala		285
	290	295
Ala Phe Asp Lys Gly Leu Glu Glu Gly Pro Asp Leu Ala Met Val Asn		300
305	310	315
Ser Ala Lys Gly Ile Thr Asn Leu His Val Pro Ser Asp Val Ile Ile		320
	325	330
Asp Ala Ser Met Pro Ala Met Ile Arg Thr Ser Gly Lys Met Trp Asn		335
	340	345
Lys Asp Asp Gln Thr Gln Asp Ala Leu Ala Val Ile Pro Asp Ser Ser		350
	355	360
Tyr Ala Gly Val Tyr Gln Thr Val Ile Glu Asp Cys Arg Lys Asn Gly		365
	370	375
Ala Phe Asp Pro Thr Thr Met Gly Thr Val Pro Asn Val Gly Leu Met		380
385	390	395
Ala Gln Lys Ala Glu Glu Tyr Gly Ser His Asp Lys Thr Phe Arg Ile		400
	405	410
Glu Ala Asp Gly Lys Val Gln Val Val Ala Ser Asn Gly Asp Val Leu		415
	420	425
Ile Glu His Asp Val Glu Lys Gly Asp Ile Trp Arg Ala Cys Gln Thr		430
	435	440
Lys Asp Ala Pro Ile Gln Asp Trp Val Lys Leu Ala Val Asn Arg Ala		445
	450	455
Arg Leu Ser Gly Met Pro Ala Val Phe Trp Leu Asp Pro Ala Arg Ala		460
465	470	475
His Asp Arg Asn Leu Thr Thr Leu Val Glu Lys Tyr Leu Ala Asp His		480
	485	490
Asp Thr Glu Gly Leu Asp Ile Gln Ile Leu Ser Pro Val Glu Ala Thr		495
	500	505
Gln His Ala Ile Asp Arg Ile Arg Arg Gly Glu Asp Thr Ile Ser Val		510
	515	520
Thr Gly Asn Val Leu Arg Asp Tyr Asn Thr Asp Leu Phe Pro Ile Leu		525
	530	535
Glu Leu Gly Thr Ser Ala Lys Met Leu Ser Val Val Pro Leu Met Ala		540
545	550	555
		560

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Gly Gly Gly Leu Phe Glu Thr Gly Ala Gly Gly Ser Ala Pro Lys His  
                             565                            570                            575  
 Val Gln Gln Val Ile Glu Glu Asn His Leu Arg Trp Asp Ser Leu Gly  
                             580                            585                            590  
 Glu Phe Leu Ala Leu Ala Glu Ser Phe Arg His Glu Leu Asn Thr Arg  
                             595                            600                            605  
 Asn Asn Thr Lys Ala Gly Val Leu Ala Asp Ala Leu Asp Arg Ala Thr  
                             610                            615                            620  
 Glu Lys Leu Leu Asn Glu Glu Lys Ser Pro Ser Arg Lys Val Gly Glu  
                             625                            630                            635                            640  
 Ile Asp Asn Arg Gly Ser His Phe Trp Leu Ala Thr Tyr Trp Ala Asp  
                             645                            650                            655  
 Glu Leu Ala Asn Gln Thr Glu Asp Ala Glu Leu Ala Glu Thr Phe Ala  
                             660                            665                            670  
 Pro Val Ala Glu Ala Leu Asn Asn Gln Ala Ala Asp Ile Asp Ala Ala  
                             675                            680                            685  
 Leu Ile Gly Glu Gln Gly Lys Pro Val Asp Leu Gly Gly Tyr Tyr Ala  
                             690                            695                            700  
 Pro Ser Asp Glu Lys Thr Ser Ala Ile Met Arg Pro Val Ala Ala Phe  
                             705                            710                            715                            720  
 Asn Glu Ile Ile Asp Ser Leu Lys Lys  
                             725

&lt;210&gt; 31

&lt;211&gt; 2322

&lt;212&gt; DNA

<213> *Corynebacterium thermoaminogenes*

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (806)..(2212)

&lt;400&gt; 31

ggtaaccccca cgtaccctag gccatcacag caattttttac atcggatatt ttaggtgtgc 60  
 tcataacgic cttatgaatt icgcagttat tagttattta aatagagaat caaacaccga 120  
 cctagccctc gccgatgcta aaagtcagct gaccccttgg ggcgcctcat ttgaaactgc 180  
 gaccaagcic atgaatgcgc gaaagcattt ccattataag ggtaagcigt aagaatagtg 240  
 ggagaaaaatg ttcatctgtg ttctaactca cttagagaaat tccatttttc tgggcctctc 300  
 tcaaatagat taagtggccc glatgctgga ttcttagaat atttagaagc gcgccaactc 360  
 atgattatgt attgtataag cctcaaagac cgaatagatt actaacattt aagttggacca 420  
 gagcgttaga agctttgtag agtgcctcat ccttgcctgac ggcaagggtt tcttaccatg 480  
 agatagatcg gcagatagtt ggttttgtaa aattttttaag gacggctcgc aatgtcaatt 540  
 ctigaacaga tcactctctt catcaacacc atcttgggtt atggctctga cgttggttct 600

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tcgccttcca	gcaacctttc	tcacacgata	ggcctgttct	aggcctaatt	ggtaalaagg	660
ctgtgttaaca	gtcgccccgcg	tgatigtgtc	tttttaggcg	cccgcgcggg	cgattttcgg	720
ttttcatctt	ttttaaatgt	agtttgggaag	atcaagtgcc	cccggatgca	cgacaatgct	780
atgccgaaca	cgtatigtig	aaatc	gtg act	gaa cat	tat gac gta gta gta	832
			Val Thr Glu His Tyr Asp	Val Val Val		
			1	5		
ctc gga gct ggc	ccc ggt ggc	tat gtc tcc	gcc atc	cgc gcc gcg	cag	880
Leu Gly Ala Gly	Pro Gly Gly	Tyr Val Ser	Ala Ile Arg	Ala Ala Gln		
10	15	20	25			
ctc ggt aag aaa	gtt gcg gtt	atc gag aag	cag tac	tgg gga ggt	gtc	928
Leu Gly Lys Lys	Val Ala Val	Ile Glu Lys	Gln Tyr Trp	Gly Gly Val		
	30	35	40			
tgc ctg aat gtg	ggt tgt atc	cca tct aag	gcg ttg	atc aag aac	gct	976
Cys Leu Asn Val	Gly Cys Ile	Pro Ser Lys	Ala Leu Ile	Lys Asn Ala		
	45	50	55			
gag atc gcc cac	atc ttc aac	cat gag aag	aag acc	ttc ggc atc	aac	1024
Glu Ile Ala His	Ile Phe Asn	His Glu Lys	Lys Thr Phe	Gly Ile Asn		
60	65	70				
ggc gag gtc acc	ttc aac tac	gag gat gcc	cac aag	cgt tcc cgt	ggt	1072
Gly Glu Val Thr	Phe Asn Tyr	Glu Asp Ala	His Lys Arg	Ser Arg Gly		
75	80	85				
gtc tcc gac aag	atc gtc ggc	ggt gtt cac	tac ttg	atg aag aag	aac	1120
Val Ser Asp Lys	Ile Val Gly	Gly Val His	Tyr Leu Met	Lys Lys Asn		
90	95	100	105			
aag atc acc gag	atc gac ggt	ttc ggc acc	ttc aag	gat gcc aag	acc	1168
Lys Ile Thr Glu	Ile Asp Gly	Phe Gly Thr	Phe Lys Asp	Ala Lys Thr		
	110	115	120			
atc gag gtg acc	gat ggt aag	gat gcc ggc	aag acc	gtc acc ttc	gat	1216
Ile Glu Val Thr	Asp Gly Lys	Asp Ala Gly	Lys Thr Val	Thr Phe Asp		
	125	130	135			
gac tgc atc atc	gcc acc ggt	tcc gtg gtc	aac tcc	ctc cgt ggt	gtt	1264
Asp Cys Ile Ile	Ala Thr Gly	Ser Val Val	Asn Ser Leu	Arg Gly Val		
	140	145	150			
gag ttc tcc gag	aac gtg gtc	tcc tac gag	gag cag	atc ctc aac	cgc	1312
Glu Phe Ser Glu	Asn Val Val	Ser Tyr Glu	Glu Gln Ile	Leu Asn Pro		
155	160	165				
gtg gcg cct aag	aag atg gtc	atc gtc ggt	ggc ggc	gcc atc ggt	atg	1360
Val Ala Pro Lys	Lys Met Val	Ile Val Gly	Gly Gly Ala	Ile Gly Met		
170	175	180	185			
gaa ttc gcc tac	gtt ctg ggc	aac tac	ggt gtg	gac gla	acc ctc atc	1408
Glu Phe Ala Tyr	Val Leu Gly	Asn Tyr Gly	Val Asp Val	Thr Leu Ile		
	190	195	200			
gag ttc atg gac	cgc gtt ctg	cgc aac	gag gat	cca gag	gtg tcc aag	1456

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Glu	Phe	Met	Asp	Arg	Val	Leu	Pro	Asn	Glu	Asp	Pro	Glu	Val	Ser	Lys	
			205					210					215			
ggt	atc	gcc	aag	gcc	tac	aag	aag	atg	ggc	atc	aag	ctc	ctc	ccg	ggc	1504
Val	Ile	Ala	Lys	Ala	Tyr	Lys	Lys	Met	Gly	Ile	Lys	Leu	Leu	Pro	Gly	
		220					225					230				
cac	gca	acc	acc	gcg	gtg	cgc	gac	aat	ggc	gat	tcc	gtt	gag	gtc	gat	1552
His	Ala	Thr	Thr	Ala	Val	Arg	Asp	Asn	Gly	Asp	Ser	Val	Glu	Val	Asp	
		235				240					245					
tac	cag	aag	aag	ggc	tcg	gac	aag	acc	gag	acc	atc	acc	gtc	gac	cgt	1600
Tyr	Gln	Lys	Lys	Gly	Ser	Asp	Lys	Thr	Glu	Thr	Ile	Thr	Val	Asp	Arg	
250					255				260					265		
gtt	ctt	atc	tcc	gtc	ggc	ttc	cgc	cca	cgc	gtc	gag	ggc	ttc	ggc	ctg	1648
Val	Leu	Ile	Ser	Val	Gly	Phe	Arg	Pro	Arg	Val	Glu	Gly	Phe	Gly	Leu	
			270					275					280			
gag	aac	acc	ggc	gtc	aag	ctc	acc	gaa	cgc	ggt	gcc	atc	gac	att	gat	1696
Glu	Asn	Thr	Gly	Val	Lys	Leu	Thr	Glu	Arg	Gly	Ala	Ile	Asp	Ile	Asp	
		285					290					295				
gag	cat	atg	cgc	acc	aac	gtc	gac	ggc	atc	tac	gcc	atc	ggt	gac	gtc	1744
Glu	His	Met	Arg	Thr	Asn	Val	Asp	Gly	Ile	Tyr	Ala	Ile	Gly	Asp	Val	
		300				305					310					
acc	gcc	aag	ctg	cag	ctg	gca	cac	gtc	gcc	gag	gca	cag	ggc	att	gtc	1792
Thr	Ala	Lys	Leu	Gln	Leu	Ala	His	Val	Ala	Glu	Ala	Gln	Gly	Ile	Val	
		315				320					325					
gcc	gcc	gag	aca	ctc	gcc	ggc	gca	gaa	acc	cag	acc	ctg	ggc	gac	tac	1840
Ala	Ala	Glu	Thr	Leu	Ala	Gly	Ala	Glu	Thr	Gln	Thr	Leu	Gly	Asp	Tyr	
330					335				340					345		
atg	atg	atg	ccg	cgt	gcc	acc	ttc	tgc	aac	cca	cag	gtt	gcc	tcc	ttc	1888
Met	Met	Met	Pro	Arg	Ala	Thr	Phe	Cys	Asn	Pro	Gln	Val	Ala	Ser	Phe	
			350					355					360			
ggt	tac	acc	gag	gag	cag	gcc	aag	gag	aag	tgg	ccg	gat	cga	gag	atc	1936
Gly	Tyr	Thr	Glu	Glu	Gln	Ala	Lys	Glu	Lys	Trp	Pro	Asp	Arg	Glu	Ile	
		365				370						375				
aag	gtg	tcc	tcc	ttc	ccg	ttc	tcc	gcg	aac	ggc	aag	gcc	gtc	ggc	ctg	1984
Lys	Val	Ser	Ser	Phe	Pro	Phe	Ser	Ala	Asn	Gly	Lys	Ala	Val	Gly	Leu	
		380				385						390				
gct	gag	acc	gat	ggt	ttc	gcc	aag	atc	gtc	gcc	gac	gct	gag	ttc	ggt	2032
Ala	Glu	Thr	Asp	Gly	Phe	Ala	Lys	Ile	Val	Ala	Asp	Ala	Glu	Phe	Gly	
		395				400					405					
gaa	ctg	ctg	ggt	ggc	cac	att	gtc	ggt	gcc	aac	gcc	tcc	gag	ctg	ctc	2080
Glu	Leu	Leu	Gly	Gly	His	Ile	Val	Gly	Ala	Asn	Ala	Ser	Glu	Leu	Leu	
410					415				420					425		
aac	gag	ctg	gtg	ctg	gcc	cag	aac	tgg	gat	ctc	acc	acc	gag	gag	atc	2128
Asn	Glu	Leu	Val	Leu	Ala	Gln	Asn	Trp	Asp	Leu	Thr	Thr	Glu	Glu	Ile	

76/123

	430	435	440	
agc cgc agc gtc cac atc cac ccg acc ctg tgc gag gct gtc aag gaa				2176
Ser Arg Ser Val His Ile His Pro Thr Leu Ser Glu Ala Val Lys Glu				
	445	450	455	
gct gcc cac ggc gtc aac ggc cac atg atc aac ttc taaatccgt				2222
Ala Ala His Gly Val Asn Gly His Met Ile Asn Phe				
	460	465		
cagacaaatg caaatccct caccgatggc atatcggiga ggggatitc tcatgcacgt				2282
aaaatcataa tccatggcaa ggaaagtcga caacagcgcc				2322

&lt;210&gt; 32

&lt;211&gt; 469

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 32

Val Thr Glu His Tyr Asp Val Val Val Leu Gly Ala Gly Pro Gly Gly				
1	5	10	15	
Tyr Val Ser Ala Ile Arg Ala Ala Gln Leu Gly Lys Lys Val Ala Val				
	20	25	30	
Ile Glu Lys Gln Tyr Trp Gly Gly Val Cys Leu Asn Val Gly Cys Ile				
	35	40	45	
Pro Ser Lys Ala Leu Ile Lys Asn Ala Glu Ile Ala His Ile Phe Asn				
	50	55	60	
His Glu Lys Lys Thr Phe Gly Ile Asn Gly Glu Val Thr Phe Asn Tyr				
	65	70	75	80
Glu Asp Ala His Lys Arg Ser Arg Gly Val Ser Asp Lys Ile Val Gly				
	85	90	95	
Gly Val His Tyr Leu Met Lys Lys Asn Lys Ile Thr Glu Ile Asp Gly				
	100	105	110	
Phe Gly Thr Phe Lys Asp Ala Lys Thr Ile Glu Val Thr Asp Gly Lys				
	115	120	125	
Asp Ala Gly Lys Thr Val Thr Phe Asp Asp Cys Ile Ile Ala Thr Gly				
	130	135	140	
Ser Val Val Asn Ser Leu Arg Gly Val Glu Phe Ser Glu Asn Val Val				
	145	150	155	160
Ser Tyr Glu Glu Gln Ile Leu Asn Pro Val Ala Pro Lys Lys Met Val				
	165	170	175	
Ile Val Gly Gly Gly Ala Ile Gly Met Glu Phe Ala Tyr Val Leu Gly				
	180	185	190	
Asn Tyr Gly Val Asp Val Thr Leu Ile Glu Phe Met Asp Arg Val Leu				
	195	200	205	
Pro Asn Glu Asp Pro Glu Val Ser Lys Val Ile Ala Lys Ala Tyr Lys				

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210	215	220
Lys Met Gly Ile Lys Leu Leu Pro Gly His Ala Thr Thr Ala Val Arg		
225	230	235
Asp Asn Gly Asp Ser Val Glu Val Asp Tyr Gln Lys Lys Gly Ser Asp		240
	245	250
Lys Thr Glu Thr Ile Thr Val Asp Arg Val Leu Ile Ser Val Gly Phe		255
	260	265
Arg Pro Arg Val Glu Gly Phe Gly Leu Glu Asn Thr Gly Val Lys Leu		270
	275	280
Thr Glu Arg Gly Ala Ile Asp Ile Asp Glu His Met Arg Thr Asn Val		285
	290	295
Asp Gly Ile Tyr Ala Ile Gly Asp Val Thr Ala Lys Leu Gln Leu Ala		300
305	310	315
His Val Ala Glu Ala Gln Gly Ile Val Ala Ala Glu Thr Leu Ala Gly		320
	325	330
Ala Glu Thr Gln Thr Leu Gly Asp Tyr Met Met Met Pro Arg Ala Thr		335
	340	345
Phe Cys Asn Pro Gln Val Ala Ser Phe Gly Tyr Thr Glu Glu Gln Ala		350
	355	360
Lys Glu Lys Trp Pro Asp Arg Glu Ile Lys Val Ser Ser Phe Pro Phe		365
	370	375
Ser Ala Asn Gly Lys Ala Val Gly Leu Ala Glu Thr Asp Gly Phe Ala		380
385	390	395
Lys Ile Val Ala Asp Ala Glu Phe Gly Glu Leu Leu Gly Gly His Ile		400
	405	410
Val Gly Ala Asn Ala Ser Glu Leu Leu Asn Glu Leu Val Leu Ala Gln		415
	420	425
Asn Trp Asp Leu Thr Thr Glu Glu Ile Ser Arg Ser Val His Ile His		430
	435	440
Pro Thr Leu Ser Glu Ala Val Lys Glu Ala Ala His Gly Val Asn Gly		445
	450	455
His Met Ile Asn Phe		460
465		

&lt;210&gt; 33

&lt;211&gt; 4096

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (250)..(3951)



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&lt;400&gt; 33

ccggatcattc gttgtttgacg ggggacgtat catcgaggat ggttcccacg atgaacttct 60  
gggagcgaat ggaacctacg caacaatgtg gcattttagta gggigacagg atatttttagg 120  
aaagacttgtt taccaaaaagg tgcataatact ggggtgtctag gtccccgcga ccggaaccag 180  
cgtttacagtg galaaaaataa agcccatlta gaaccttcaa caagcaagga aaagaggcga 240  
gtaccigcc gtg agc agc gct agt act ttc ggc cag aac gcg tgg ctg gtg 291

Val Ser Ser Ala Ser Thr Phe Gly Gln Asn Ala Trp Leu Val

10

gat gag atg ttc cag cag ttc aag aag gac ccc cag tcc gtg gac aag 339  
Asp Glu Met Phe Gln Gln Phe Lys Lys Asp Pro Gln Ser Val Asp Lys

30

gaa tgg aga gag ctc ttc gag tct cag ggg ggt ccc cag gct gaa aag 387  
Glu Trp Arg Glu Leu Phe Glu Ser Gln Gly Gly Pro Gln Ala Glu Lys

45

gct acc ccc gcc acc ccc gaa gcc aag aag gca gct tcg tcg cag tcc 435  
Ala Thr Pro Ala Thr Pro Glu Ala Lys Lys Ala Ala Ser Ser Gln Ser

60

tca act tcc gga cag tcc acc gcc aag gct gcc cct gcc gcc aag acc 483  
Ser Thr Ser Gly Gln Ser Thr Ala Lys Ala Ala Pro Ala Ala Lys Thr

75

gca ccg gcc tci gcg cca gcc aag gcl gcc ccl gtl aag caa aac cag 531  
Ala Pro Ala Ser Ala Pro Ala Lys Ala Ala Pro Val Lys Gln Asn Gln

90

gcg tcc aag ccl gcc aag aag gcc aag gag tcc ccc ctg tcc aag cca 579  
Ala Ser Lys Pro Ala Lys Lys Ala Lys Glu Ser Pro Leu Ser Lys Pro

11

gct gcc atg cct gag ccg gga acc acc cca ctc agg ggc atc ttc aag 627  
Ala Ala Met Pro Glu Pro Gly Thr Thr Pro Leu Arg Gly Ile Phe Lys

1.25

tcc atc gcc aag aac atg gac ctc tcc ctc gag gtg ccc acc gcc acc 675  
 Ser Ile Ala Lys Asn Met Asp Leu Ser Leu Glu Val Pro Thr Ala Thr

140

tcc gtc cgc gac atg ccc gcg cgc ctc atg ttc gag aac cgc gcc atg 723  
 Ser Val Arg Asp Met Pro Ala Arg Leu Met Phe Glu Asn Arg Ala Met

155

gtc aac gac cag ctc aag cgc acc cgt ggc ggc aag atc tcc ttc acc 771  
Val Asn Asp Gln Leu Lys Arg Thr Arg Gly Gly Lys Ile Ser Phe Thr

170

cac atc atc ggc tac gcc atg gtg aag gct gtc atg gca cac ccg gac 819  
His Ile Ile Gly Tyr Ala Met Val Lys Ala Val Met Ala His Pro Asp

175                      180                      185                      190  
atg aac aac tcc tat gac atc gtc gac ggc aag ccg tcc ctg gtc gtc 867

Met Asn Asn Ser Tyr Asp Ile Val Asp Gly Lys Pro Ser Leu Val Val

79/123

			195				200				205						
ccg	gag	cac	atc	aac	ctc	ggc	ctg	gcc	atc	gac	ctc	ccc	cag	aag	gac	915	
Pro	Glu	His	Ile	Asn	Leu	Gly	Leu	Ala	Ile	Asp	Leu	Pro	Gln	Lys	Asp		
			210				215				220						
ggc	tcc	cgt	gcc	ctc	gtg	gtc	gcc	gcc	atc	aag	gaa	acc	gag	aag	atg	963	
Gly	Ser	Arg	Ala	Leu	Val	Val	Ala	Ala	Ile	Lys	Glu	Thr	Glu	Lys	Met		
			225				230				235						
acc	ttc	tcc	cag	ttc	ctg	gag	gcc	tat	gag	gac	gtt	gtg	gca	cgc	tcc	1011	
Thr	Phe	Ser	Gln	Phe	Leu	Glu	Ala	Tyr	Glu	Asp	Val	Val	Ala	Arg	Ser		
			240				245				250						
cgc	gtc	ggc	aag	ctc	acc	atg	gat	gac	tac	cag	ggt	gtc	acc	atc	tcc	1059	
Arg	Val	Gly	Lys	Leu	Thr	Met	Asp	Asp	Tyr	Gln	Gly	Val	Thr	Ile	Ser		
255			260				265				270						
tig	acc	aac	ccg	ggt	ggc	atc	ggt	acc	cgc	cac	tcc	atc	ccg	cgt	ctg	1107	
Leu	Thr	Asn	Pro	Gly	Gly	Ile	Gly	Thr	Arg	His	Ser	Ile	Pro	Arg	Leu		
			275				280				285						
acc	aag	ggc	cag	ggc	acc	atc	atc	ggt	gtc	ggt	tcc	atg	gac	tac	ccg	1155	
Thr	Lys	Gly	Gln	Gly	Thr	Ile	Ile	Gly	Val	Gly	Ser	Met	Asp	Tyr	Pro		
			290				295				300						
gcc	gag	ttc	cag	ggt	gcc	tcc	gag	gac	cgt	ctc	gcc	gag	ctc	ggt	gtg	1203	
Ala	Glu	Phe	Gln	Gly	Ala	Ser	Glu	Asp	Arg	Leu	Ala	Glu	Leu	Gly	Val		
			305				310				315						
ggc	aag	ctc	gtc	acc	atc	acc	tcc	acc	tac	gat	cac	cgc	gtc	atc	cag	1251	
Gly	Lys	Leu	Val	Thr	Ile	Thr	Ser	Thr	Tyr	Asp	His	Arg	Val	Ile	Gln		
			320				325				330						
ggc	gcg	gaa	tcc	ggt	gag	ttc	ctg	cgc	acc	atg	tcc	cag	ctg	ctc	gtg	1299	
Gly	Ala	Glu	Ser	Gly	Glu	Phe	Leu	Arg	Thr	Met	Ser	Gln	Leu	Leu	Val		
335			340				345				350						
gac	gat	gcg	ttc	tgg	gat	cac	atc	ttc	gag	gag	atg	aac	gtt	ccc	tac	1347	
Asp	Asp	Ala	Phe	Trp	Asp	His	Ile	Phe	Glu	Glu	Met	Asn	Val	Pro	Tyr		
			355				360				365						
acc	ccg	atg	cgc	tgg	gca	cag	gac	ctg	ccc	aac	acc	ggt	gtg	gac	aag	1395	
Thr	Pro	Met	Arg	Trp	Ala	Gln	Asp	Leu	Pro	Asn	Thr	Gly	Val	Asp	Lys		
			370				375				380						
aac	acc	cgt	gtc	atg	cag	ctc	atc	gag	gcc	tac	cgc	tcc	cgc	ggt	cac	1443	
Asn	Thr	Arg	Val	Met	Gln	Leu	Ile	Glu	Ala	Tyr	Arg	Ser	Arg	Gly	His		
			385				390				395						
ctc	atc	gcc	gac	acc	aac	cca	ctg	ccc	tgg	gtc	cag	ccc	ggc	atg	ccc	1491	
Leu	Ile	Ala	Asp	Thr	Asn	Pro	Leu	Pro	Trp	Val	Gln	Pro	Gly	Met	Pro		
			400				405				410						
gtc	ccg	gat	cac	cgt	gac	ctc	gac	atc	gag	acc	cac	ggc	ctg	acc	ctg	1539	
Val	Pro	Asp	His	Arg	Asp	Leu	Asp	Ile	Glu	Thr	His	Gly	Leu	Thr	Leu		
415			420				425				430						

80/123

tgg gat ctg gac cgt acc ttc cac gtc ggt ggt ttc ggt ggc aag gag	1587
Trp Asp Leu Asp Arg Thr Phe His Val Gly Gly Phe Gly Gly Lys Glu	
435 440 445	
acc atg acc ctg cgc gag gtg ctc agc cgc ctc cgc gcc gcc tac acc	1635
Thr Met Thr Leu Arg Glu Val Leu Ser Arg Leu Arg Ala Ala Tyr Thr	
450 455 460	
ctc aag gtc ggc tcc gag tac acc cac atc ctc gac cgc gat gag cgc	1683
Leu Lys Val Gly Ser Glu Tyr Thr His Ile Leu Asp Arg Asp Glu Arg	
465 470 475	
acc tgg ctg cag gac cgc ctc gag gcc ggt atg ccc aag ccc acc gcc	1731
Thr Trp Leu Gln Asp Arg Leu Glu Ala Gly Met Pro Lys Pro Thr Ala	
480 485 490	
gcc gag cag aag tac atc ctg cag aag ctc aac gcc gcc gag gca ttc	1779
Ala Glu Gln Lys Tyr Ile Leu Gln Lys Leu Asn Ala Ala Glu Ala Phe	
495 500 505 510	
gag aac ttc ctg cag acc aag tac gtc ggc cag aag cgt ttc tcc ctc	1827
Glu Asn Phe Leu Gln Thr Lys Tyr Val Gly Gln Lys Arg Phe Ser Leu	
515 520 525	
gag ggt gcc gag tca ctg atc ccg ctg atg gac tcc gcc atc gac acc	1875
Glu Gly Ala Glu Ser Leu Ile Pro Leu Met Asp Ser Ala Ile Asp Thr	
530 535 540	
gcc gca ggc cag ggc ctt gac gag gtc gtc atc ggc atg ccc cac cgt	1923
Ala Ala Gly Gln Gly Leu Asp Glu Val Val Ile Gly Met Pro His Arg	
545 550 555	
ggt cgc ctc aac gtg ctg ttc aac atc gtc ggc aag cca ctg gcc tcc	1971
Gly Arg Leu Asn Val Leu Phe Asn Ile Val Gly Lys Pro Leu Ala Ser	
560 565 570	
atc ttc aac gag ttc gag ggc cag atg gag cag ggc cag atc ggt ggc	2019
Ile Phe Asn Glu Phe Glu Gly Gln Met Glu Gln Gly Gln Ile Gly Gly	
575 580 585 590	
tcc ggt gac gtg aag tac cac ctc ggt tcc gag ggc acc cac ctg cag	2067
Ser Gly Asp Val Lys Tyr His Leu Gly Ser Glu Gly Thr His Leu Gln	
595 600 605	
atg ttc ggc gac ggc gag atc aag gtc tcc ctc acc gcc aac ccc tcc	2115
Met Phe Gly Asp Gly Glu Ile Lys Val Ser Leu Thr Ala Asn Pro Ser	
610 615 620	
cac ctc gag gcc gtc aac ccg gtc gtg gag ggc atc gtc cgc gcc aag	2163
His Leu Glu Ala Val Asn Pro Val Val Glu Gly Ile Val Arg Ala Lys	
625 630 635	
cag gac atc ctg gac aag ggc ccg gac ggc tac acc gtc gtc ccg ctg	2211
Gln Asp Ile Leu Asp Lys Gly Pro Asp Gly Tyr Thr Val Val Pro Leu	
640 645 650	
ctg ctc cac ggt gac gcc gcc ttc gcc ggc ctg ggc atc gtg ccc gag	2259

81/123

Leu	Leu	His	Gly	Asp	Ala	Ala	Phe	Ala	Gly	Leu	Gly	Ile	Val	Pro	Glu	
655					660					665					670	
acc	atc	aac	ctc	gca	gcc	ctg	cgt	ggt	tac	gat	gtc	ggt	ggc	acc	atc	2307
Thr	Ile	Asn	Leu	Ala	Ala	Leu	Arg	Gly	Tyr	Asp	Val	Gly	Gly	Thr	Ile	
			675						680					685		
cac	atc	gtg	gtc	aac	aac	cag	atc	ggc	ttc	acc	acc	acc	ccg	gac	tcc	2355
His	Ile	Val	Val	Asn	Asn	Gln	Ile	Gly	Phe	Thr	Thr	Thr	Pro	Asp	Ser	
			690					695					700			
agc	cgt	tcc	atg	cac	tac	gcc	acc	gac	tgc	gcc	aag	gcc	ttc	ggt	tgc	2403
Ser	Arg	Ser	Met	His	Tyr	Ala	Thr	Asp	Cys	Ala	Lys	Ala	Phe	Gly	Cys	
		705					710					715				
ccg	gtg	ttc	cac	gtc	aac	ggt	gac	gac	ccc	gag	gct	gig	gtc	tgg	gtc	2451
Pro	Val	Phe	His	Val	Asn	Gly	Asp	Asp	Pro	Glu	Ala	Val	Val	Trp	Val	
	720					725					730					
ggc	cag	ctg	gcc	acc	gag	tac	cgt	cgc	cgc	ttc	ggc	aag	gat	gtc	ttc	2499
Gly	Gln	Leu	Ala	Thr	Glu	Tyr	Arg	Arg	Arg	Phe	Gly	Lys	Asp	Val	Phe	
735					740					745				750		
atc	gac	ctc	atc	tgc	tac	cgc	ctg	cgc	ggc	cac	aac	gag	gct	gat	gac	2547
Ile	Asp	Leu	Ile	Cys	Tyr	Arg	Leu	Arg	Gly	His	Asn	Glu	Ala	Asp	Asp	
				755					760					765		
cca	tcc	atg	acc	cag	ccg	aag	atg	tac	gag	ctg	atc	acc	ggc	cgc	gac	2595
Pro	Ser	Met	Thr	Gln	Pro	Lys	Met	Tyr	Glu	Leu	Ile	Thr	Gly	Arg	Asp	
			770					775					780			
tcc	gtg	cgt	gcc	acc	tac	acc	gag	gac	ctc	ctc	ggc	cgt	ggt	gac	ctc	2643
Ser	Val	Arg	Ala	Thr	Tyr	Thr	Glu	Asp	Leu	Leu	Gly	Arg	Gly	Asp	Leu	
		785					790					795				
tcc	ccc	gag	gac	gcc	gag	gcc	gtt	gtc	cgc	gac	ttc	cac	gac	cag	atg	2691
Ser	Pro	Glu	Asp	Ala	Glu	Ala	Val	Val	Arg	Asp	Phe	His	Asp	Gln	Met	
	800					805					810					
gaa	tcc	gtg	ttc	aac	gag	gtc	aag	gaa	gcc	ggc	aag	aag	cag	cct	gat	2739
Glu	Ser	Val	Phe	Asn	Glu	Val	Lys	Glu	Ala	Gly	Lys	Lys	Gln	Pro	Asp	
815					820					825				830		
gag	cag	acc	ggc	atc	acc	ggt	tcc	cag	gaa	ctg	acc	cgt	ggc	ctg	gac	2787
Glu	Gln	Thr	Gly	Ile	Thr	Gly	Ser	Gln	Glu	Leu	Thr	Arg	Gly	Leu	Asp	
			835						840				845			
acc	aac	atc	acc	cgc	gag	gaa	ctg	gtc	gaa	ctc	ggc	cag	gcc	ttc	gtc	2835
Thr	Asn	Ile	Thr	Arg	Glu	Glu	Leu	Val	Glu	Leu	Gly	Gln	Ala	Phe	Val	
			850					855					860			
aac	acc	cca	gag	ggc	ttc	acc	tac	cac	cca	cgt	gtg	gca	ccg	gtg	gcc	2883
Asn	Thr	Pro	Glu	Gly	Phe	Thr	Tyr	His	Pro	Arg	Val	Ala	Pro	Val	Ala	
		865					870					875				
aag	aag	cgt	gcc	gag	tcc	gtc	acc	gag	ggt	ggc	atc	gac	tgg	gca	tgg	2931
Lys	Lys	Arg	Ala	Glu	Ser	Val	Thr	Glu	Gly	Gly	Ile	Asp	Trp	Ala	Trp	

82/123

880	885	890	
ggc gag ctc atc gcc ttc ggc tcc ctg gcc acc tcc ggc agg ctg gtc			2979
Gly Glu Leu Ile Ala Phe Gly Ser Leu Ala Thr Ser Gly Arg Leu Val			
895	900	905	910
cgc ctc gcc ggt gag gat tcc cgc cgt ggt acc ttc acc cag cgt cac			3027
Arg Leu Ala Gly Glu Asp Ser Arg Arg Gly Thr Phe Thr Gln Arg His			
	915	920	925
gcc gtg gcc atc gac ccg aac acc gcc gag gag ttc aac ccg ctc cac			3075
Ala Val Ala Ile Asp Pro Asn Thr Ala Glu Glu Phe Asn Pro Leu His			
	930	935	940
gag ctg gca cag gcc aag ggc ggc ggc aag ttc ctc gtc tac aac tcc			3123
Glu Leu Ala Gln Ala Lys Gly Gly Gly Lys Phe Leu Val Tyr Asn Ser			
	945	950	955
gcg ctg acc gag tac gcg ggt atg ggc ttc gaa tac ggc tac tcc gtg			3171
Ala Leu Thr Glu Tyr Ala Gly Met Gly Phe Glu Tyr Gly Tyr Ser Val			
	960	965	970
ggc aac ccg gac gcc gtg gtg tcc tgg gag gca cag ttc ggt gac ttc			3219
Gly Asn Pro Asp Ala Val Val Ser Trp Glu Ala Gln Phe Gly Asp Phe			
	975	980	985
ggc aac ggt gca cag acc atc atc gat gag tac atc tcc tcc ggt gag			3267
Ala Asn Gly Ala Gln Thr Ile Ile Asp Glu Tyr Ile Ser Ser Gly Glu			
	995	1000	1005
gcc aag tgg ggc cag acc tcc tcc gtc atc ctg ctg ctg ccc cac ggt			3315
Ala Lys Trp Gly Gln Thr Ser Ser Val Ile Leu Leu Leu Pro His Gly			
	1010	1015	1020
tac gag ggc cag ggt ccg gac cac tcc tcc gca cgc atc gag cgt ttc			3363
Tyr Glu Gly Gln Gly Pro Asp His Ser Ser Ala Arg Ile Glu Arg Phe			
	1025	1030	1035
ctg cag ctg tgc gcc gag ggt tcc atg acc atc gcc cag ccg acc acc			3411
Leu Gln Leu Cys Ala Glu Gly Ser Met Thr Ile Ala Gln Pro Thr Thr			
	1040	1045	1050
ccg gcg aac tac ttc cac ctg ctg cgt cgt cac gca ctg ggc aag atg			3459
Pro Ala Asn Tyr Phe His Leu Leu Arg Arg His Ala Leu Gly Lys Met			
	1055	1060	1065
aag cgc ccg ctg gtc gtc ttc acc ccg aag tcc atg ctg cgc aac aag			3507
Lys Arg Pro Leu Val Val Phe Thr Pro Lys Ser Met Leu Arg Asn Lys			
	1075	1080	1085
gcc gcc acc tcc gct ccg gag gag ttc acc gag gtc acc cgc ttc aag			3555
Ala Ala Thr Ser Ala Pro Glu Glu Phe Thr Glu Val Thr Arg Phe Lys			
	1090	1095	1100
tcc gtg atc gac gat ccg aac gtg gcg gat gcc tcc aag gtg aag aag			3603
Ser Val Ile Asp Asp Pro Asn Val Ala Asp Ala Ser Lys Val Lys Lys			
	1105	1110	1115

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```

atc atg ctg tgc tcc ggc aag atc tac tac gaa ctg gcc aag cgc aag 3651
Ile Met Leu Cys Ser Gly Lys Ile Tyr Tyr Glu Leu Ala Lys Arg Lys
1120 1125 1130
gag aag gac aac cgc gac gac atc gcg atc gtg cgc atc gag atg ctg 3699
Glu Lys Asp Asn Arg Asp Asp Ile Ala Ile Val Arg Ile Glu Met Leu
1135 1140 1145 1150
cac ccg atc ccg ttc aac cgt ctg cgc gac gcc ttc gac ggc tac ccc 3747
His Pro Ile Pro Phe Asn Arg Leu Arg Asp Ala Phe Asp Gly Tyr Pro
1155 1160 1165
aac gcc gag gag atc ctg ttc gtt cag gac gag ccg gca aac cag ggt 3795
Asn Ala Glu Glu Ile Leu Phe Val Gln Asp Glu Pro Ala Asn Gln Gly
1170 1175 1180
gcc tgg ccg ttc tac cag gag cac ctg ccc aac ctg atc gag ggc atg 3843
Ala Trp Pro Phe Tyr Gln Glu His Leu Pro Asn Leu Ile Glu Gly Met
1185 1190 1195
ctc ccg atg cgt cgc atc tgc cgc cgt tcc cag tcc tgc act gcg acc 3891
Leu Pro Met Arg Arg Ile Ser Arg Arg Ser Gln Ser Ser Thr Ala Thr
1200 1205 1210
ggt atc gcg aag gtg cac acc atc gag cag cag aag ctg ctg gat gat 3939
Gly Ile Ala Lys Val His Thr Ile Glu Gln Gln Lys Leu Leu Asp Asp
1215 1220 1225 1230
gcg ttc aac gca taaacgttaa tacagcgggt gatacctga accccgccgc 3991
Ala Phe Asn Ala
accttllaga tgcgggcggg gttllgctll gcctgcatag gcgataatat tcatatacac 4051
ccatcacgtt taagttctgc atttggatcg tgcgagcatc ccggt 4096

```

&lt;210&gt; 34

&lt;211&gt; 1234

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 34

```

Val Ser Ser Ala Ser Thr Phe Gly Gln Asn Ala Trp Leu Val Asp Glu
1 5 10 15
Met Phe Gln Gln Phe Lys Lys Asp Pro Gln Ser Val Asp Lys Glu Trp
20 25 30
Arg Glu Leu Phe Glu Ser Gln Gly Gly Pro Gln Ala Glu Lys Ala Thr
35 40 45
Pro Ala Thr Pro Glu Ala Lys Lys Ala Ala Ser Ser Gln Ser Ser Thr
50 55 60
Ser Gly Gln Ser Thr Ala Lys Ala Ala Pro Ala Ala Lys Thr Ala Pro
65 70 75 80
Ala Ser Ala Pro Ala Lys Ala Ala Pro Val Lys Gln Asn Gln Ala Ser

```

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85												90				95			
Lys	Pro	Ala	Lys	Lys	Ala	Lys	Glu	Ser	Pro	Leu	Ser	Lys	Pro	Ala	Ala				
100						105						110							
Met	Pro	Glu	Pro	Gly	Thr	Thr	Pro	Leu	Arg	Gly	Ile	Phe	Lys	Ser	Ile				
115						120						125							
Ala	Lys	Asn	Met	Asp	Leu	Ser	Leu	Glu	Val	Pro	Thr	Ala	Thr	Ser	Val				
130						135						140							
Arg	Asp	Met	Pro	Ala	Arg	Leu	Met	Phe	Glu	Asn	Arg	Ala	Met	Val	Asn				
145						150						155							
Asp	Gln	Leu	Lys	Arg	Thr	Arg	Gly	Gly	Lys	Ile	Ser	Phe	Thr	His	Ile				
			165						170										
Ile	Gly	Tyr	Ala	Met	Val	Lys	Ala	Val	Met	Ala	His	Pro	Asp	Met	Asn				
			180						185			190							
Asn	Ser	Tyr	Asp	Ile	Val	Asp	Gly	Lys	Pro	Ser	Leu	Val	Val	Pro	Glu				
195						200						205							
His	Ile	Asn	Leu	Gly	Leu	Ala	Ile	Asp	Leu	Pro	Gln	Lys	Asp	Gly	Ser				
210						215						220							
Arg	Ala	Leu	Val	Val	Ala	Ala	Ile	Lys	Glu	Thr	Glu	Lys	Met	Thr	Phe				
225						230						235							
Ser	Gln	Phe	Leu	Glu	Ala	Tyr	Glu	Asp	Val	Val	Ala	Arg	Ser	Arg	Val				
			245						250										
Gly	Lys	Leu	Thr	Met	Asp	Asp	Tyr	Gln	Gly	Val	Thr	Ile	Ser	Leu	Thr				
			260						265			270							
Asn	Pro	Gly	Gly	Ile	Gly	Thr	Arg	His	Ser	Ile	Pro	Arg	Leu	Thr	Lys				
275						280						285							
Gly	Gln	Gly	Thr	Ile	Ile	Gly	Val	Gly	Ser	Met	Asp	Tyr	Pro	Ala	Glu				
290						295						300							
Phe	Gln	Gly	Ala	Ser	Glu	Asp	Arg	Leu	Ala	Glu	Leu	Gly	Val	Gly	Lys				
305						310						315							
Leu	Val	Thr	Ile	Thr	Ser	Thr	Tyr	Asp	His	Arg	Val	Ile	Gln	Gly	Ala				
			325						330										
Glu	Ser	Gly	Glu	Phe	Leu	Arg	Thr	Met	Ser	Gln	Leu	Leu	Val	Asp	Asp				
			340						345			350							
Ala	Phe	Trp	Asp	His	Ile	Phe	Glu	Glu	Met	Asn	Val	Pro	Tyr	Thr	Pro				
355						360						365							
Met	Arg	Trp	Ala	Gln	Asp	Leu	Pro	Asn	Thr	Gly	Val	Asp	Lys	Asn	Thr				
370						375						380							
Arg	Val	Met	Gln	Leu	Ile	Glu	Ala	Tyr	Arg	Ser	Arg	Gly	His	Leu	Ile				
385						390						395							
Ala	Asp	Thr	Asn	Pro	Leu	Pro	Trp	Val	Gln	Pro	Gly	Met	Pro	Val	Pro				
			405						410										
Asp	His	Arg	Asp	Leu	Asp	Ile	Glu	Thr	His	Gly	Leu	Thr	Leu	Trp	Asp				
			420						425			430							

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Leu	Asp	Arg	Thr	Phe	His	Val	Gly	Gly	Phe	Gly	Gly	Lys	Glu	Thr	Met
		435					440					445			
Thr	Leu	Arg	Glu	Val	Leu	Ser	Arg	Leu	Arg	Ala	Ala	Tyr	Thr	Leu	Lys
	450					455					460				
Val	Gly	Ser	Glu	Tyr	Thr	His	Ile	Leu	Asp	Arg	Asp	Glu	Arg	Thr	Trp
465					470				475						480
Leu	Gln	Asp	Arg	Leu	Glu	Ala	Gly	Met	Pro	Lys	Pro	Thr	Ala	Ala	Glu
			485					490					495		
Gln	Lys	Tyr	Ile	Leu	Gln	Lys	Leu	Asn	Ala	Ala	Glu	Ala	Phe	Glu	Asn
			500				505						510		
Phe	Leu	Gln	Thr	Lys	Tyr	Val	Gly	Gln	Lys	Arg	Phe	Ser	Leu	Glu	Gly
	515					520					525				
Ala	Glu	Ser	Leu	Ile	Pro	Leu	Met	Asp	Ser	Ala	Ile	Asp	Thr	Ala	Ala
	530					535				540					
Gly	Gln	Gly	Leu	Asp	Glu	Val	Val	Ile	Gly	Met	Pro	His	Arg	Gly	Arg
545					550				555						560
Leu	Asn	Val	Leu	Phe	Asn	Ile	Val	Gly	Lys	Pro	Leu	Ala	Ser	Ile	Phe
			565				570						575		
Asn	Glu	Phe	Glu	Gly	Gln	Met	Glu	Gln	Gly	Gln	Ile	Gly	Gly	Ser	Gly
			580				585					590			
Asp	Val	Lys	Tyr	His	Leu	Gly	Ser	Glu	Gly	Thr	His	Leu	Gln	Met	Phe
	595					600						605			
Gly	Asp	Gly	Glu	Ile	Lys	Val	Ser	Leu	Thr	Ala	Asn	Pro	Ser	His	Leu
	610					615				620					
Glu	Ala	Val	Asn	Pro	Val	Val	Glu	Gly	Ile	Val	Arg	Ala	Lys	Gln	Asp
625					630				635						640
Ile	Leu	Asp	Lys	Gly	Pro	Asp	Gly	Tyr	Thr	Val	Val	Pro	Leu	Leu	Leu
			645				650					655			
His	Gly	Asp	Ala	Ala	Phe	Ala	Gly	Leu	Gly	Ile	Val	Pro	Glu	Thr	Ile
			660				665					670			
Asn	Leu	Ala	Ala	Leu	Arg	Gly	Tyr	Asp	Val	Gly	Gly	Thr	Ile	His	Ile
	675					680						685			
Val	Val	Asn	Asn	Gln	Ile	Gly	Phe	Thr	Thr	Thr	Pro	Asp	Ser	Ser	Arg
	690					695				700					
Ser	Met	His	Tyr	Ala	Thr	Asp	Cys	Ala	Lys	Ala	Phe	Gly	Cys	Pro	Val
705					710				715						720
Phe	His	Val	Asn	Gly	Asp	Asp	Pro	Glu	Ala	Val	Val	Trp	Val	Gly	Gln
			725				730					735			
Leu	Ala	Thr	Glu	Tyr	Arg	Arg	Arg	Phe	Gly	Lys	Asp	Val	Phe	Ile	Asp
		740					745					750			
Leu	Ile	Cys	Tyr	Arg	Leu	Arg	Gly	His	Asn	Glu	Ala	Asp	Asp	Pro	Ser
	755					760					765				
Met	Thr	Gln	Pro	Lys	Met	Tyr	Glu	Leu	Ile	Thr	Gly	Arg	Asp	Ser	Val



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770		775		780
Arg Ala Thr Tyr Thr Glu Asp Leu Leu Gly Arg Gly Asp Leu Ser Pro				
785		790		795
Glu Asp Ala Glu Ala Val Val Arg Asp Phe His Asp Gln Met Glu Ser				800
		805		810
Val Phe Asn Glu Val Lys Glu Ala Gly Lys Lys Gln Pro Asp Glu Gln				815
		820		825
Thr Gly Ile Thr Gly Ser Gln Glu Leu Thr Arg Gly Leu Asp Thr Asn				830
		835		840
Ile Thr Arg Glu Glu Leu Val Glu Leu Gly Gln Ala Phe Val Asn Thr				845
		850		855
Pro Glu Gly Phe Thr Tyr His Pro Arg Val Ala Pro Val Ala Lys Lys				860
865		870		875
Arg Ala Glu Ser Val Thr Glu Gly Gly Ile Asp Trp Ala Trp Gly Glu				880
		885		890
Leu Ile Ala Phe Gly Ser Leu Ala Thr Ser Gly Arg Leu Val Arg Leu				895
		900		905
Ala Gly Glu Asp Ser Arg Arg Gly Thr Phe Thr Gln Arg His Ala Val				910
		915		920
Ala Ile Asp Pro Asn Thr Ala Glu Glu Phe Asn Pro Leu His Glu Leu				925
		930		935
Ala Gln Ala Lys Gly Gly Gly Lys Phe Leu Val Tyr Asn Ser Ala Leu				940
945		950		955
Thr Glu Tyr Ala Gly Met Gly Phe Glu Tyr Gly Tyr Ser Val Gly Asn				960
		965		970
Pro Asp Ala Val Val Ser Trp Glu Ala Gln Phe Gly Asp Phe Ala Asn				975
		980		985
Gly Ala Gln Thr Ile Ile Asp Glu Tyr Ile Ser Ser Gly Glu Ala Lys				990
		995		1000
Trp Gly Gln Thr Ser Ser Val Ile Leu Leu Leu Pro His Gly Tyr Glu				1005
1010		1015		1020
Gly Gln Gly Pro Asp His Ser Ser Ala Arg Ile Glu Arg Phe Leu Gln				1025
		1030		1035
Leu Cys Ala Glu Gly Ser Met Thr Ile Ala Gln Pro Thr Thr Pro Ala				1040
		1045		1050
Asn Tyr Phe His Leu Leu Arg Arg His Ala Leu Gly Lys Met Lys Arg				1055
		1060		1065
Pro Leu Val Val Phe Thr Pro Lys Ser Met Leu Arg Asn Lys Ala Ala				1070
		1075		1080
Thr Ser Ala Pro Glu Glu Phe Thr Glu Val Thr Arg Phe Lys Ser Val				1085
		1090		1095
Ile Asp Asp Pro Asn Val Ala Asp Ala Ser Lys Val Lys Lys Ile Met				1100
1105		1110		1115
				1120

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Leu Cys Ser Gly Lys Ile Tyr Tyr Glu Leu Ala Lys Arg Lys Glu Lys  
                     1125                    1130                    1135  
 Asp Asn Arg Asp Asp Ile Ala Ile Val Arg Ile Glu Met Leu His Pro  
                     1140                    1145                    1150  
 Ile Pro Phe Asn Arg Leu Arg Asp Ala Phe Asp Gly Tyr Pro Asn Ala  
                     1155                    1160                    1165  
 Glu Glu Ile Leu Phe Val Gln Asp Glu Pro Ala Asn Gln Gly Ala Trp  
                     1170                    1175                    1180  
 Pro Phe Tyr Gln Glu His Leu Pro Asn Leu Ile Glu Gly Met Leu Pro  
 185                    1190                    1195                    1200  
 Met Arg Arg Ile Ser Arg Arg Ser Gln Ser Ser Thr Ala Thr Gly Ile  
                     1205                    1210                    1215  
 Ala Lys Val His Thr Ile Glu Gln Gln Lys Leu Leu Asp Asp Ala Phe  
                     1220                    1225                    1230  
 Asn Ala

&lt;210&gt; 35

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for aceA

&lt;400&gt; 35

ccctctaccca gcgaactccg

20

&lt;210&gt; 36

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for aceA

&lt;400&gt; 36

ctgccttgaa ctacagggttc

20

&lt;210&gt; 37

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

88/123

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for accBC

&lt;400&gt; 37

catccacccc ggctacggct

20

&lt;210&gt; 38

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for accBC

&lt;400&gt; 38

cggtgactgg glgttcacc

20

&lt;210&gt; 39

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for dtsR1

&lt;400&gt; 39

acggcccagc cctgaccgac

20

&lt;210&gt; 40

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for dtsR1

&lt;400&gt; 40

agcagcgccc atgacggcga

20

&lt;210&gt; 41

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

89/123

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for dtsR2

&lt;400&gt; 41

acggcccagc cctgaccgac

20

&lt;210&gt; 42

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for dtsR2

&lt;400&gt; 42

agcagcgccc atgacggcga

20

&lt;210&gt; 43

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for pfk

&lt;400&gt; 43

cgtcatccga ggaatcgtcc

20

&lt;210&gt; 44

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for pfk

&lt;400&gt; 44

cgtggcggcc catgacctcc

21

&lt;210&gt; 45

&lt;211&gt; 17

&lt;212&gt; DNA

$\langle 220 \rangle$  $\langle 220 \rangle$ 

<221> UNSURE

$$\langle 222 \rangle \quad (3)$$

$\langle 223 \rangle$  n=a or g or c or t

<400> 45

ggncghytba aygaycc

17

$\langle 210 \rangle$  46

211 20

## <212> DNA

### 〈213〉 Artificial Sequence

 $\langle 220 \rangle$ 

<223> Description of Artificial Sequence: primer for scrB

 $\langle 220 \rangle$ 

&lt;221&gt; UNSURE

 $\langle 222 \rangle$  (18)

$\langle 223 \rangle$  n=a or g or c or t

<400> 46

ggrcaylccc acatrtancc

20

<210> 47

 $\langle 211 \rangle$  20

## <212> DNA

### <213> Artificial Sequence

 $\langle 220 \rangle$ 

⟨223⟩ Description of Artificial Sequence: primer for gluABCD

<400> 47

ccatccggat ccggcaagtc

20

$\langle 210 \rangle$  48

 $\langle 211 \rangle$  20

## <212> DNA

91/123

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: primer for gluABCD

<400> 48

aatcccatct cgtgggtaac

20

<210> 49

<211> 23

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: primer for pdhA

<400> 49

actgtgtcca tgggtcttgg ccc

23

<210> 50

<211> 20

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: primer for pdhA

<400> 50

cgctggaatccgaacatcga

20

<210> 51

<211> 26

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: primer for pc

<400> 51

ggcgcaacct acgacgttgc aatgcg

26

<210> 52

<211> 20

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&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for pc

&lt;400&gt; 52

tggccgcctg ggatctcgtg

20

&lt;210&gt; 53

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for ppc

&lt;400&gt; 53

ggttccctgga ttggtggaga

20

&lt;210&gt; 54

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for ppc

&lt;400&gt; 54

ccgccatcct tgttggaatc

20

&lt;210&gt; 55

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for acn

&lt;220&gt;

&lt;221&gt; UNSURE

&lt;222&gt; (3, 6, 9)

&lt;223&gt; n=inosine

93/123

<400> 55  
ginggnacng aytcsatac 20

<210> 56  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for acn

<220>  
<221> UNSURE  
<222> (3,9,18)  
<223> n=inosine

<400> 56  
gcnggagana tgtgrtcngt 20

<210> 57  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for icd

<400> 57  
gacatttcac tcgctggacg 20

<210> 58  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for icd

<400> 58  
ccgtactctt cagccttctg 20

<210> 59



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&lt;211&gt; 17

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for lpd

&lt;400&gt; 59

atcatcgcaa ccggttc

17

&lt;210&gt; 60

&lt;211&gt; 19

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for lpd

&lt;400&gt; 60

cgtcaccgat ggcgtaaat

19

&lt;210&gt; 61

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for odhA

&lt;400&gt; 61

acaccgtggt cgccccaacg

20

&lt;210&gt; 62

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for odhA

&lt;400&gt; 62

tgctaaccgg tcccacctgg

20

95/123

&lt;210&gt; 63

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
screening PCR of lpd

&lt;400&gt; 63

tacgaggagc agalcctcaa

20

&lt;210&gt; 64

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
screening PCR of lpd

&lt;400&gt; 64

ttgacgccgg tgttctccag

20

&lt;210&gt; 65

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
LA cloning of acn

&lt;400&gt; 65

ggtgaagcta agtagttagc

20

&lt;210&gt; 66

&lt;211&gt; 18

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence: primer for

96/123

LA cloning of acn

<400> 66  
agctactaaa cctgcacc 18

<210> 67  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of icd

<400> 67  
ccgtactctt cagccttcig 67

<210> 68  
<211> 18  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of icd

<400> 68  
tcgtccttgt tccacatc 18

<210> 69  
<211> 17  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of lpd

<400> 69  
atcatcgcaa ccggttc 17

<210> 70  
<211> 20

97/123

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
LA cloning of lpd

&lt;400&gt; 70

tacgaggagc agatccctcaa

20

&lt;210&gt; 71

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
LA cloning of acn

&lt;400&gt; 71

gctaacctact tagcttcacc

20

&lt;210&gt; 72

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
LA cloning of acn

&lt;400&gt; 72

gaaccaggaa ctattgaacc

20

&lt;210&gt; 73

&lt;211&gt; 18

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
LA cloning of icd

98/123

<400> 73  
tccgatgtca tcatcgac 18

<210> 74  
<211> 18  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of icd

<400> 74  
atgtggaaca aggacgac 18

<210> 75  
<211> 35  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of odhA

<400> 75  
gtacatatlg tcgttagaac gcgtaatacg actca 35

<210> 76  
<211> 35  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: primer for  
LA cloning of odhA

<400> 76  
cgttagaacg cgtaatacga ctcactatag ggaga 35

<210> 77  
<211> 32  
<212> DNA

99/123

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gdh gene

&lt;400&gt; 77

gcgcctgcag gtccgagggg gtgcgttcgg ca

32

&lt;210&gt; 78

&lt;211&gt; 32

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gdh gene

&lt;400&gt; 78

gcgcctgcag ccaccagga tgcctcaacc ag

32

&lt;210&gt; 79

&lt;211&gt; 1344

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1)..(1341)

&lt;400&gt; 79

atg act gta gat gag cag gtc tcc aac tac tac gac atg ctg ctg aag 48  
Met Thr Val Asp Glu Gln Val Ser Asn Tyr Tyr Asp Met Leu Leu Lys

1

5

10

15

cgc aac gcc ggg gaa cct gag ttc cac cag gct gtc gcg gag gtt ctc 96  
Arg Asn Ala Gly Glu Pro Glu Phe His Gln Ala Val Ala Glu Val Leu

20

25

30

gaa tct ctg aag atc gtc ctg gag aag gac ccg cac tac gcc gac tac 144  
Glu Ser Leu Lys Ile Val Leu Glu Lys Asp Pro His Tyr Ala Asp Tyr

35

40

45

ggi ctg atc cag cgt ctc tgc gaa ccg gaa cgc cag ctg atc ttc cgt 192

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Gly	Leu	Ile	Gln	Arg	Leu	Cys	Glu	Pro	Glu	Arg	Gln	Leu	Ile	Phe	Arg		
50						55					60						
gtg	ccc	igg	gtg	gat	gac	aac	ggt	cag	gtg	cac	gtc	aac	cgt	ggt	ttc	240	
Val	Pro	Trp	Val	Asp	Asp	Asn	Gly	Gln	Val	His	Val	Asn	Arg	Gly	Phe		
65						70					75				80		
cgt	gtc	cag	ttc	aac	tcc	gca	ctc	ggc	ccg	tac	aag	ggt	ggt	ctg	cgt	288	
Arg	Val	Gln	Phe	Asn	Ser	Ala	Leu	Gly	Pro	Tyr	Lys	Gly	Gly	Leu	Arg		
				85					90					95			
ttc	cac	ccc	tcc	gtc	aac	ctc	ggc	atc	gtc	aag	ttc	ctc	ggc	ttc	gag	336	
Phe	His	Pro	Ser	Val	Asn	Leu	Gly	Ile	Val	Lys	Phe	Leu	Gly	Phe	Glu		
			100					105				110					
cag	atc	ttc	aag	aac	tcc	ctc	acc	ggt	ctg	ccg	atc	ggt	ggc	ggc	aag	384	
Gln	Ile	Phe	Lys	Asn	Ser	Leu	Thr	Gly	Leu	Pro	Ile	Gly	Gly	Gly	Lys		
		115				120					125						
ggt	ggt	tcc	gac	ttc	gac	ccg	aag	ggc	aag	tcc	gag	ctg	gag	atc	atg	432	
Gly	Gly	Ser	Asp	Phe	Asp	Pro	Lys	Gly	Lys	Ser	Glu	Leu	Glu	Ile	Met		
	130					135					140						
cgc	ttc	tgc	cag	tcc	ttc	atg	acc	gag	ctg	cac	cgc	cac	atc	ggc	gag	480	
Arg	Phe	Cys	Gln	Ser	Phe	Met	Thr	Glu	Leu	His	Arg	His	Ile	Gly	Glu		
145					150					155				160			
tac	cgg	gat	gtc	ccg	gcc	ggt	gac	atc	gga	gtc	ggt	ggc	cgc	gag	atc	528	
Tyr	Arg	Asp	Val	Pro	Ala	Gly	Asp	Ile	Gly	Val	Gly	Gly	Arg	Glu	Ile		
			165					170					175				
ggt	tac	ctc	ttc	ggc	cac	tac	cgc	cgt	ctg	gcc	aac	cag	cac	gag	tcc	576	
Gly	Tyr	Leu	Phe	Gly	His	Tyr	Arg	Arg	Leu	Ala	Asn	Gln	His	Glu	Ser		
		180					185				190						
ggt	gtg	ctc	acc	ggc	aag	ggc	ctg	acc	tgg	ggt	ggt	tcc	ctg	gtc	cgc	624	
Gly	Val	Leu	Thr	Gly	Lys	Gly	Leu	Thr	Trp	Gly	Gly	Ser	Leu	Val	Arg		
	195					200					205						
acc	gag	gcc	acc	ggc	ttc	ggc	acc	gtc	tac	ttc	gtc	cag	gag	atg	atc	672	
Thr	Glu	Ala	Thr	Gly	Phe	Gly	Thr	Val	Tyr	Phe	Val	Gln	Glu	Met	Ile		
	210					215					220						
aag	gcg	gaa	ggg	gag	acc	ctc	gag	ggc	aag	aag	gtc	atc	gtc	tcc	ggt	720	
Lys	Ala	Glu	Gly	Glu	Thr	Leu	Glu	Gly	Lys	Lys	Val	Ile	Val	Ser	Gly		
225					230					235				240			
tcc	ggc	aac	gtg	gcc	acc	tac	gcc	atc	cag	aag	gtg	cag	gaa	ctg	ggt	768	
Ser	Gly	Asn	Val	Ala	Thr	Tyr	Ala	Ile	Gln	Lys	Val	Gln	Glu	Leu	Gly		
		245					250					255					
gcg	gtt	gtg	gtc	ggc	ttc	tcc	gac	tcc	agc	ggc	tgg	gtc	tcc	acc	ccg	816	
Ala	Val	Val	Val	Gly	Phe	Ser	Asp	Ser	Ser	Gly	Trp	Val	Ser	Thr	Pro		
		260					265				270						
aac	ggt	gtt	gac	gtg	gcc	aag	ctg	cgt	gag	atc	aag	gag	gtc	cgt	cgt	864	
Asn	Gly	Val	Asp	Val	Ala	Lys	Leu	Arg	Glu	Ile	Lys	Glu	Val	Arg	Arg		

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275	280	285	
gca cgc gtg tcc tcc tac gcc gac gag gtg gag ggt gcg gag tac cac			912
Ala Arg Val Ser Ser Tyr Ala Asp Glu Val Glu Gly Ala Glu Tyr His			
290	295	300	
acc gac ggc tcc atc tgg gat ctg acc gcc gac atc gcg ctg ccc tgc			960
Thr Asp Gly Ser Ile Trp Asp Leu Thr Ala Asp Ile Ala Leu Pro Cys			
305	310	315	320
gcc acc cag aac gaa ctg gac ggc gac aac gcc cgc acc ctc gcg gac			1008
Ala Thr Gln Asn Glu Leu Asp Gly Asp Asn Ala Arg Thr Leu Ala Asp			
325	330	335	
aac ggc tgc cgc ttc gtg gcg gag ggc gcc aac atg ccc tcc acc ccc			1056
Asn Gly Cys Arg Phe Val Ala Glu Gly Ala Asn Met Pro Ser Thr Pro			
340	345	350	
gag gcc atc gac gtc ttc cgt gag cgt ggt gtt ctc ttc ggg ccg ggc			1104
Glu Ala Ile Asp Val Phe Arg Glu Arg Gly Val Leu Phe Gly Pro Gly			
355	360	365	
aag gct gcc aac gcc ggt ggc gtg gcc acc tcc gcc ctg gag atg cag			1152
Lys Ala Ala Asn Ala Gly Gly Val Ala Thr Ser Ala Leu Glu Met Gln			
370	375	380	
cag aac gcc tcc cgt gat tcc tgg agc ttc gag tac acc gat gag cgt			1200
Gln Asn Ala Ser Arg Asp Ser Trp Ser Phe Glu Tyr Thr Asp Glu Arg			
385	390	395	400
ctc cac cgc atc atg aag aac atc ttc aag tcc tgc gcc gat acc gcc			1248
Leu His Arg Ile Met Lys Asn Ile Phe Lys Ser Cys Ala Asp Thr Ala			
405	410	415	
aag gag tac ggc cac gag aag aac tac gtg gtc ggt gcg aac atc gcc			1296
Lys Glu Tyr Gly His Glu Lys Asn Tyr Val Val Gly Ala Asn Ile Ala			
420	425	430	
gga ttc aag aag gtc gct gac gcc atg ctc gcc cag ggt gtc atc taa			1344
Gly Phe Lys Lys Val Ala Asp Ala Met Leu Ala Gln Gly Val Ile			
435	440	445	

&lt;210&gt; 80

&lt;211&gt; 447

&lt;212&gt; PRT

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;400&gt; 80

Met Thr Val Asp Glu Gln Val Ser Asn Tyr Tyr Asp Met Leu Leu Lys	
1 5 10 15	
Arg Asn Ala Gly Glu Pro Glu Phe His Gln Ala Val Ala Glu Val Leu	
20 25 30	
Glu Ser Leu Lys Ile Val Leu Glu Lys Asp Pro His Tyr Ala Asp Tyr	



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35					40					45						
Gly	Leu	Ile	Gln	Arg	Leu	Cys	Glu	Pro	Glu	Arg	Gln	Leu	Ile	Phe	Arg	
50					55					60						
Val	Pro	Trp	Val	Asp	Asp	Asn	Gly	Gln	Val	His	Val	Asn	Arg	Gly	Phe	
65					70					75					80	
Arg	Val	Gln	Phe	Asn	Ser	Ala	Leu	Gly	Pro	Tyr	Lys	Gly	Gly	Leu	Arg	
85					90					95						
Phe	His	Pro	Ser	Val	Asn	Leu	Gly	Ile	Val	Lys	Phe	Leu	Gly	Phe	Glu	
100					105					110						
Gln	Ile	Phe	Lys	Asn	Ser	Leu	Thr	Gly	Leu	Pro	Ile	Gly	Gly	Gly	Lys	
115					120					125						
Gly	Gly	Ser	Asp	Phe	Asp	Pro	Lys	Gly	Lys	Ser	Glu	Leu	Glu	Ile	Met	
130					135					140						
Arg	Phe	Cys	Gln	Ser	Phe	Met	Thr	Glu	Leu	His	Arg	His	Ile	Gly	Glu	
145					150					155					160	
Tyr	Arg	Asp	Val	Pro	Ala	Gly	Asp	Ile	Gly	Val	Gly	Gly	Arg	Glu	Ile	
165					170					175						
Gly	Tyr	Leu	Phe	Gly	His	Tyr	Arg	Arg	Leu	Ala	Asn	Gln	His	Glu	Ser	
180					185					190						
Gly	Val	Leu	Thr	Gly	Lys	Gly	Leu	Thr	Trp	Gly	Gly	Ser	Leu	Val	Arg	
195					200					205						
Thr	Glu	Ala	Thr	Gly	Phe	Gly	Thr	Val	Tyr	Phe	Val	Gln	Glu	Met	Ile	
210					215					220						
Lys	Ala	Glu	Gly	Glu	Thr	Leu	Glu	Gly	Lys	Lys	Val	Ile	Val	Ser	Gly	
225					230					235					240	
Ser	Gly	Asn	Val	Ala	Thr	Tyr	Ala	Ile	Gln	Lys	Val	Gln	Glu	Leu	Gly	
245					250					255						
Ala	Val	Val	Val	Gly	Phe	Ser	Asp	Ser	Ser	Gly	Trp	Val	Ser	Thr	Pro	
260					265					270						
Asn	Gly	Val	Asp	Val	Ala	Lys	Leu	Arg	Glu	Ile	Lys	Glu	Val	Arg	Arg	
275					280					285						
Ala	Arg	Val	Ser	Ser	Tyr	Ala	Asp	Glu	Val	Glu	Gly	Ala	Glu	Tyr	His	
290					295					300						
Thr	Asp	Gly	Ser	Ile	Trp	Asp	Leu	Thr	Ala	Asp	Ile	Ala	Leu	Pro	Cys	
305					310					315					320	
Ala	Thr	Gln	Asn	Glu	Leu	Asp	Gly	Asp	Asn	Ala	Arg	Thr	Leu	Ala	Asp	
325					330					335						
Asn	Gly	Cys	Arg	Phe	Val	Ala	Glu	Gly	Ala	Asn	Met	Pro	Ser	Thr	Pro	
340					345					350						
Glu	Ala	Ile	Asp	Val	Phe	Arg	Glu	Arg	Gly	Val	Leu	Phe	Gly	Pro	Gly	
355					360					365						
Lys	Ala	Ala	Asn	Ala	Gly	Gly	Val	Ala	Thr	Ser	Ala	Leu	Glu	Met	Gln	
370					375					380						

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Gln Asn Ala Ser Arg Asp Ser Trp Ser Phe Glu Tyr Thr Asp Glu Arg  
 385 390 395 400  
 Leu His Arg Ile Met Lys Asn Ile Phe Lys Ser Cys Ala Asp Thr Ala  
 405 410 415  
 Lys Glu Tyr Gly His Glu Lys Asn Tyr Val Val Gly Ala Asn Ile Ala  
 420 425 430  
 Gly Phe Lys Lys Val Ala Asp Ala Met Leu Ala Gln Gly Val Ile  
 435 440 445

&lt;210&gt; 81

&lt;211&gt; 1344

&lt;212&gt; DNA

&lt;213&gt; Brevibacterium lactofermentum

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1).. (1341)

&lt;400&gt; 81

atg aca gtt gat gag cag gtc tct aac tat tac gac atg ctt ctg aag 48  
 Met Thr Val Asp Glu Gln Val Ser Asn Tyr Tyr Asp Met Leu Leu Lys  
 1 5 10 15  
 cgc aat gct ggc gag cct gaa ttt cac cag gca gtg gca gag gtt ttg 96  
 Arg Asn Ala Gly Glu Pro Glu Phe His Gln Ala Val Ala Glu Val Leu  
 20 25 30  
 gaa tct ttg aag atc gtc ctg gaa aag gac cct cat tac gct gat tac 144  
 Glu Ser Leu Lys Ile Val Leu Glu Lys Asp Pro His Tyr Ala Asp Tyr  
 35 40 45  
 ggt ctc atc cag cgc ctg tgc gag cct gag cgt cag ctc atc ttc cgt 192  
 Gly Leu Ile Gln Arg Leu Cys Glu Pro Glu Arg Gln Leu Ile Phe Arg  
 50 55 60  
 gtg cct tgg gtt gat gac cag ggc cag gtc cac gtc aac cgt ggt ttc 240  
 Val Pro Trp Val Asp Asp Gln Gly Gln Val His Val Asn Arg Gly Phe  
 65 70 75 80  
 cgc gtg cag ttc aac tct gca ctt gga cca tac aag ggc ggc ctg cgc 288  
 Arg Val Gln Phe Asn Ser Ala Leu Gly Pro Tyr Lys Gly Gly Leu Arg  
 85 90 95  
 ttc cac cca tct gta aac ctg ggc att gtg aag ttc ctg ggc ttt gag 336  
 Phe His Pro Ser Val Asn Leu Gly Ile Val Lys Phe Leu Gly Phe Glu  
 100 105 110  
 cag atc ttt aaa aac tcc cta acc ggc ctg cca atc ggt ggt ggc aag 384  
 Gln Ile Phe Lys Asn Ser Leu Thr Gly Leu Pro Ile Gly Gly Gly Lys  
 115 120 125

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ggt gga tcc gac ttc gac cct aag ggc aag tcc gat ctg gaa atc atg	432
Gly Gly Ser Asp Phe Asp Pro Lys Gly Lys Ser Asp Leu Glu Ile Met	
130 135 140	
cgt ttc tgc cag tcc ttc atg acc gag ctg cac cgc cac atc ggt gag	480
Arg Phe Cys Gln Ser Phe Met Thr Glu Leu His Arg His Ile Gly Glu	
145 150 155 160	
tac cgc gac gtt cct gca ggt gac atc gga gtt ggt ggc cgc gag atc	528
Tyr Arg Asp Val Pro Ala Gly Asp Ile Gly Val Gly Gly Arg Glu Ile	
165 170 175	
ggt tac ctg ttt ggc cac tac cgt cgc atg gct aac cag cac gag tcc	576
Gly Tyr Leu Phe Gly His Tyr Arg Arg Met Ala Asn Gln His Glu Ser	
180 185 190	
ggc gtt ttg acc ggt aag ggc ctg acc tgg ggt gga tcc ctg gtc cgc	624
Gly Val Leu Thr Gly Lys Gly Leu Thr Trp Gly Gly Ser Leu Val Arg	
195 200 205	
acc gag gca act ggc tac ggc tgc gtt tac ttc gtg agt gaa atg atc	672
Thr Glu Ala Thr Gly Tyr Gly Cys Val Tyr Phe Val Ser Glu Met Ile	
210 215 220	
aag gct aag ggc gag agc atc agc ggc cag aag atc atc gtt tcc ggt	720
Lys Ala Lys Gly Glu Ser Ile Ser Gly Gln Lys Ile Ile Val Ser Gly	
225 230 235 240	
tcc ggc aac gta gca acc tac gcg att gaa aag gct cag gaa ctc ggc	768
Ser Gly Asn Val Ala Thr Tyr Ala Ile Glu Lys Ala Gln Glu Leu Gly	
245 250 255	
gca acc gtt att ggt ttc tcc gat tcc agc ggt tgg gtt cat acc cct	816
Ala Thr Val Ile Gly Phe Ser Asp Ser Ser Gly Trp Val His Thr Pro	
260 265 270	
aac ggc gtt gac gtg gct aag ctg cgc gaa atc aag gaa gtt cgc cgc	864
Asn Gly Val Asp Val Ala Lys Leu Arg Glu Ile Lys Glu Val Arg Arg	
275 280 285	
gca cgc gta tcc gtg tac gcc gac gaa att gaa ggc gca acc tac cac	912
Ala Arg Val Ser Val Tyr Ala Asp Glu Ile Glu Gly Ala Thr Tyr His	
290 295 300	
acc gac ggt tcc atc tgg gat ctg aag tgc gat atc gct ctt cct tgt	960
Thr Asp Gly Ser Ile Trp Asp Leu Lys Cys Asp Ile Ala Leu Pro Cys	
305 310 315 320	
gca act cag aac gag ctg aac ggc gag aac gct aag act ctt gca gag	1008
Ala Thr Gln Asn Glu Leu Asn Gly Glu Asn Ala Lys Thr Leu Ala Asp	
325 330 335	
aac ggc tgc cgt ttc gtt gct gaa ggc gcg aac atg cct tcc acc cct	1056
Asn Gly Cys Arg Phe Val Ala Glu Gly Ala Asn Met Pro Ser Thr Pro	
340 345 350	
gag gct gtt gag gtc ttc cgt gag cgc gac atc cgc ttc gga cca ggc	1104

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Glu	Ala	Val	Glu	Val	Phe	Arg	Glu	Arg	Asp	Ile	Arg	Phe	Gly	Pro	Gly		
		355						360					365				
aag	gca	gct	aac	gct	ggt	ggc	gtt	gca	acc	tcc	gct	ctg	gag	atg	cag	1152	
Lys	Ala	Ala	Asn	Ala	Gly	Gly	Val	Ala	Thr	Ser	Ala	Leu	Glu	Met	Gln		
		370				375					380						
cag	aac	gct	tcg	cgc	gat	tcc	tgg	agc	ttc	gag	tac	acc	gac	gag	cgc	1200	
Gln	Asn	Ala	Ser	Arg	Asp	Ser	Trp	Ser	Phe	Glu	Tyr	Thr	Asp	Glu	Arg		
		385			390					395					400		
ctc	cag	gtg	atc	atg	aag	aac	atc	ttc	aag	acc	tgt	gca	gag	acc	gca	1248	
Leu	Gln	Val	Ile	Met	Lys	Asn	Ile	Phe	Lys	Thr	Cys	Ala	Glu	Thr	Ala		
			405						410					415			
gca	gag	tat	gga	cac	gag	aac	gat	tac	gtt	gtc	ggc	gct	aac	att	gct	1296	
Ala	Glu	Tyr	Gly	His	Glu	Asn	Asp	Tyr	Val	Val	Gly	Ala	Asn	Ile	Ala		
		420					425					430					
ggc	ttt	aag	aag	gta	gct	gac	gcg	atg	ctg	gca	cag	ggc	gtc	atc	taa	1344	
Gly	Phe	Lys	Lys	Val	Ala	Asp	Ala	Met	Leu	Ala	Gln	Gly	Val	Ile			
		435					440					445					

&lt;210&gt; 82

&lt;211&gt; 447

&lt;212&gt; PRT

&lt;213&gt; Brevibacterium lactofermentum

&lt;400&gt; 82

Met	Thr	Val	Asp	Glu	Gln	Val	Ser	Asn	Tyr	Tyr	Asp	Met	Leu	Leu	Lys		
1				5					10					15			
Arg	Asn	Ala	Gly	Glu	Pro	Glu	Phe	His	Gln	Ala	Val	Ala	Glu	Val	Leu		
		20					25						30				
Glu	Ser	Leu	Lys	Ile	Val	Leu	Glu	Lys	Asp	Pro	His	Tyr	Ala	Asp	Tyr		
		35				40						45					
Gly	Leu	Ile	Gln	Arg	Leu	Cys	Glu	Pro	Glu	Arg	Gln	Leu	Ile	Phe	Arg		
	50				55					60							
Val	Pro	Trp	Val	Asp	Asp	Gln	Gly	Gln	Val	His	Val	Asn	Arg	Gly	Phe		
	65			70					75					80			
Arg	Val	Gln	Phe	Asn	Ser	Ala	Leu	Gly	Pro	Tyr	Lys	Gly	Gly	Leu	Arg		
		85					90						95				
Phe	His	Pro	Ser	Val	Asn	Leu	Gly	Ile	Val	Lys	Phe	Leu	Gly	Phe	Glu		
		100				105						110					
Gln	Ile	Phe	Lys	Asn	Ser	Leu	Thr	Gly	Leu	Pro	Ile	Gly	Gly	Gly	Lys		
	115					120						125					
Gly	Gly	Ser	Asp	Phe	Asp	Pro	Lys	Gly	Lys	Ser	Asp	Leu	Glu	Ile	Met		
	130				135					140							
Arg	Phe	Cys	Gln	Ser	Phe	Met	Thr	Glu	Leu	His	Arg	His	Ile	Gly	Glu		

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145		150		155		160									
Tyr	Arg	Asp	Val	Pro	Ala	Gly	Asp	Ile	Gly	Val	Gly	Gly	Arg	Glu	Ile
				165					170					175	
Gly	Tyr	Leu	Phe	Gly	His	Tyr	Arg	Arg	Met	Ala	Asn	Gln	His	Glu	Ser
			180					185					190		
Gly	Val	Leu	Thr	Gly	Lys	Gly	Leu	Thr	Trp	Gly	Gly	Ser	Leu	Val	Arg
		195					200					205			
Thr	Glu	Ala	Thr	Gly	Tyr	Gly	Cys	Val	Tyr	Phe	Val	Ser	Glu	Met	Ile
	210					215				220					
Lys	Ala	Lys	Gly	Glu	Ser	Ile	Ser	Gly	Gln	Lys	Ile	Ile	Val	Ser	Gly
225					230					235					240
Ser	Gly	Asn	Val	Ala	Thr	Tyr	Ala	Ile	Glu	Lys	Ala	Gln	Glu	Leu	Gly
			245					250					255		
Ala	Thr	Val	Ile	Gly	Phe	Ser	Asp	Ser	Ser	Gly	Trp	Val	His	Thr	Pro
		260					265					270			
Asn	Gly	Val	Asp	Val	Ala	Lys	Leu	Arg	Glu	Ile	Lys	Glu	Val	Arg	Arg
	275					280				285					
Ala	Arg	Val	Ser	Val	Tyr	Ala	Asp	Glu	Ile	Glu	Gly	Ala	Thr	Tyr	His
	290				295					300					
Thr	Asp	Gly	Ser	Ile	Trp	Asp	Leu	Lys	Cys	Asp	Ile	Ala	Leu	Pro	Cys
305				310					315						320
Ala	Thr	Gln	Asn	Glu	Leu	Asn	Gly	Glu	Asn	Ala	Lys	Thr	Leu	Ala	Asp
			325				330						335		
Asn	Gly	Cys	Arg	Phe	Val	Ala	Glu	Gly	Ala	Asn	Met	Pro	Ser	Thr	Pro
		340					345					350			
Glu	Ala	Val	Glu	Val	Phe	Arg	Glu	Arg	Asp	Ile	Arg	Phe	Gly	Pro	Gly
	355					360				365					
Lys	Ala	Ala	Asn	Ala	Gly	Gly	Val	Ala	Thr	Ser	Ala	Leu	Glu	Met	Gln
	370				375				380						
Gln	Asn	Ala	Ser	Arg	Asp	Ser	Trp	Ser	Phe	Glu	Tyr	Thr	Asp	Glu	Arg
385			390						395						400
Leu	Gln	Val	Ile	Met	Lys	Asn	Ile	Phe	Lys	Thr	Cys	Ala	Glu	Thr	Ala
			405				410					415			
Ala	Glu	Tyr	Gly	His	Glu	Asn	Asp	Tyr	Val	Val	Gly	Ala	Asn	Ile	Ala
		420					425					430			
Gly	Phe	Lys	Lys	Val	Ala	Asp	Ala	Met	Leu	Ala	Gln	Gly	Val	Ile	
	435					440					445				

&lt;210&gt; 83

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

107/123

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;220&gt;

&lt;221&gt; misc\_feature

&lt;222&gt; (9)

&lt;223&gt; n=inosine

&lt;400&gt; 83

aagatcacnt acatcgaygg

20

&lt;210&gt; 84

&lt;211&gt; 20

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 84

tagaagtcta cgttcgggta

20

&lt;210&gt; 85

&lt;211&gt; 21

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 85

gtcgacaata gcctgaatct g

21

&lt;210&gt; 86

&lt;211&gt; 21

&lt;212&gt; DNA

108/123

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 86

cggtggaacc ggtgctgaca t

21

&lt;210&gt; 87

&lt;211&gt; 21

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 87

ggglgggga attcggtcatt t

21°

&lt;210&gt; 88

&lt;211&gt; 21

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 88

tgicgtagcc gcggtagcgc a

21

&lt;210&gt; 89

&lt;211&gt; 1293

&lt;212&gt; DNA

&lt;213&gt; Corynebacterium thermoaminogenes

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1)..(1290)

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&lt;400&gt; 89

gtg gct tct gat aac aac aag gct gta ctg cac tac cct ggc ggc gaa	48
Val Ala Ser Asp Asn Asn Lys Ala Val Leu His Tyr Pro Gly Gly Glu	
1 5 10 15	
ttc gag atg ggc atc aag cag gcc acc gag ggt aac tcc ggt gtc atc	96
Phe Glu Met Gly Ile Lys Gln Ala Thr Glu Gly Asn Ser Gly Val Ile	
20 25 30	
ctg ggt aag atg ctg tcg gaa acc ggt ctg gtc acc ttc gac ccc ggt	144
Leu Gly Lys Met Leu Ser Glu Thr Gly Leu Val Thr Phe Asp Pro Gly	
35 40 45	
tat gtc agc acc ggt tcc acc gaa tcc aag atc acc tac atc gat ggt	192
Tyr Val Ser Thr Gly Ser Thr Glu Ser Lys Ile Thr Tyr Ile Asp Gly	
50 55 60	
gat gca ggc atc ctg cgc tac cgc ggc tac gac att gcg gat ctg gcc	240
Asp Ala Gly Ile Leu Arg Tyr Arg Gly Tyr Asp Ile Ala Asp Leu Ala	
65 70 75 80	
gaa aat gcc acc ttc aat gag gtc tcc tac ctc ctg atc aag ggt gag	288
Glu Asn Ala Thr Phe Asn Glu Val Ser Tyr Leu Leu Ile Lys Gly Glu	
85 90 95	
ctc ccg acc ccg gaa gag ctc cac aag ttc aac gac gag att cgt cac	336
Leu Pro Thr Pro Glu Glu Leu His Lys Phe Asn Asp Glu Ile Arg His	
100 105 110	
cac acc ctg ctg gac gag gac ttc aag tcc cag ttc aat gtc ttc cct	384
His Thr Leu Leu Asp Glu Asp Phe Lys Ser Gln Phe Asn Val Phe Pro	
115 120 125	
cgc gat gcc cac ccg atg gcc acc ctg gcc tcc tcg gtt aac atc ctc	432
Arg Asp Ala His Pro Met Ala Thr Leu Ala Ser Ser Val Asn Ile Leu	
130 135 140	
tcc acc tac tac cag gat cag ctg gat ccc ctg gat gag gct cag ctg	480
Ser Thr Tyr Tyr Gln Asp Gln Leu Asp Pro Leu Asp Glu Ala Gln Leu	
145 150 155 160	
gac aag gca acc gtc cgc ctg atg gcg aag gtt ccg atg ctg gct gca	528
Asp Lys Ala Thr Val Arg Leu Met Ala Lys Val Pro Met Leu Ala Ala	
165 170 175	
tac gca cac cgt gcc cgc aag ggt gcg ccg tac atg tac ccg gac aac	576
Tyr Ala His Arg Ala Arg Lys Gly Ala Pro Tyr Met Tyr Pro Asp Asn	
180 185 190	
tcc ctc aat gcc cgt gag aac ttc ctg cgc atg atg ttc ggt tac ccg	624
Ser Leu Asn Ala Arg Glu Asn Phe Leu Arg Met Met Phe Gly Tyr Pro	
195 200 205	
acc gag ccg tac gag gtt gat ccg atc atg gtc aaa gcc ctc gac aag	672
Thr Glu Pro Tyr Glu Val Asp Pro Ile Met Val Lys Ala Leu Asp Lys	



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210	215	220	
ctg ctc atc ctg cac gca gac cac gag cag aac tgc tcc acc tcc act			720
Leu Leu Ile Leu His Ala Asp His Glu Gln Asn Cys Ser Thr Ser Thr			
225	230	235	240
gtc cgc atg atc ggc tcc gcg cag gcg aac atg ttc gtc tcc atc gcc			768
Val Arg Met Ile Gly Ser Ala Gln Ala Asn Met Phe Val Ser Ile Ala			
	245	250	255
ggc ggc atc aac gca ctc tcc ggc ccg ctg cac ggt ggc gcc aac cag			816
Gly Gly Ile Asn Ala Leu Ser Gly Pro Leu His Gly Gly Ala Asn Gln			
	260	265	270
gct gtc ctc gag atg ctc gag gag atc gca gcc aac ggc ggc gac gca			864
Ala Val Leu Glu Met Leu Glu Glu Ile Ala Ala Asn Gly Gly Asp Ala			
	275	280	285
acc gac ttc atg aac cgc gtg aag aac aag gag aag ggt gtc cgc ctc			912
Thr Asp Phe Met Asn Arg Val Lys Asn Lys Glu Lys Gly Val Arg Leu			
	290	295	300
atg ggc ttc gga cac cgc gtc tac aag aac tac gat ccg cgt gca gcc			960
Met Gly Phe Gly His Arg Val Tyr Lys Asn Tyr Asp Pro Arg Ala Ala			
	305	310	315
atc gtc aag gac acc gcc cac gag atc ctc gag cac ctc ggt ggc gac			1008
Ile Val Lys Asp Thr Ala His Glu Ile Leu Glu His Leu Gly Gly Asp			
	325	330	335
cca ctg ctg gat ctg gct ctc aag ctg gaa gaa atc gca ctc aac gac			1056
Pro Leu Leu Asp Leu Ala Leu Lys Leu Glu Glu Ile Ala Leu Asn Asp			
	340	345	350
gat tac ttc atc tcc cgc aag ctg tac ccg aac gtg gac ttc tac acc			1104
Asp Tyr Phe Ile Ser Arg Lys Leu Tyr Pro Asn Val Asp Phe Tyr Thr			
	355	360	365
ggc ctg atc tac cgc gcc atg ggc ttc ccg acg gac ttc ttc acc gtc			1152
Gly Leu Ile Tyr Arg Ala Met Gly Phe Pro Thr Asp Phe Phe Thr Val			
	370	375	380
ctg ttc gcc atc ggc cgc ctc ccg ggc tgg atc gcc cac tac cgc gag			1200
Leu Phe Ala Ile Gly Arg Leu Pro Gly Trp Ile Ala His Tyr Arg Glu			
	385	390	395
cag ctc gcc gat ccg ggc gcc aag atc aac cgt cct cgc cag atc tac			1248
Gln Leu Ala Asp Pro Gly Ala Lys Ile Asn Arg Pro Arg Gln Ile Tyr			
	405	410	415
acc ggt gag acc gca cgc aag atc atc ccc cgc gaa gag cgc tag			1293
Thr Gly Glu Thr Ala Arg Lys Ile Ile Pro Arg Glu Glu Arg			
	420	425	430

&lt;210&gt; 90

&lt;211&gt; 430

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&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 90

Val	Ala	Ser	Asp	Asn	Asn	Lys	Ala	Val	Leu	His	Tyr	Pro	Gly	Gly	Glu
1				5				10						15	
Phe	Glu	Met	Gly	Ile	Lys	Gln	Ala	Thr	Glu	Gly	Asn	Ser	Gly	Val	Ile
			20					25					30		
Leu	Gly	Lys	Met	Leu	Ser	Glu	Thr	Gly	Leu	Val	Thr	Phe	Asp	Pro	Gly
		35					40					45			
Tyr	Val	Ser	Thr	Gly	Ser	Thr	Glu	Ser	Lys	Ile	Thr	Tyr	Ile	Asp	Gly
	50					55					60				
Asp	Ala	Gly	Ile	Leu	Arg	Tyr	Arg	Gly	Tyr	Asp	Ile	Ala	Asp	Leu	Ala
65					70					75				80	
Glu	Asn	Ala	Thr	Phe	Asn	Glu	Val	Ser	Tyr	Leu	Leu	Ile	Lys	Gly	Glu
				85						90				95	
Leu	Pro	Thr	Pro	Glu	Glu	Leu	His	Lys	Phe	Asn	Asp	Glu	Ile	Arg	His
			100					105					110		
His	Thr	Leu	Leu	Asp	Glu	Asp	Phe	Lys	Ser	Gln	Phe	Asn	Val	Phe	Pro
		115					120					125			
Arg	Asp	Ala	His	Pro	Met	Ala	Thr	Leu	Ala	Ser	Ser	Val	Asn	Ile	Leu
	130					135					140				
Ser	Thr	Tyr	Tyr	Gln	Asp	Gln	Leu	Asp	Pro	Leu	Asp	Glu	Ala	Gln	Leu
145				150					155					160	
Asp	Lys	Ala	Thr	Val	Arg	Leu	Met	Ala	Lys	Val	Pro	Met	Leu	Ala	Ala
			165						170				175		
Tyr	Ala	His	Arg	Ala	Arg	Lys	Gly	Ala	Pro	Tyr	Met	Tyr	Pro	Asp	Asn
		180					185					190			
Ser	Leu	Asn	Ala	Arg	Glu	Asn	Phe	Leu	Arg	Met	Met	Phe	Gly	Tyr	Pro
	195					200					205				
Thr	Glu	Pro	Tyr	Glu	Val	Asp	Pro	Ile	Met	Val	Lys	Ala	Leu	Asp	Lys
	210					215					220				
Leu	Leu	Ile	Leu	His	Ala	Asp	His	Glu	Gln	Asn	Cys	Ser	Thr	Ser	Thr
225				230					235					240	
Val	Arg	Met	Ile	Gly	Ser	Ala	Gln	Ala	Asn	Met	Phe	Val	Ser	Ile	Ala
			245						250				255		
Gly	Gly	Ile	Asn	Ala	Leu	Ser	Gly	Pro	Leu	His	Gly	Gly	Ala	Asn	Gln
		260					265					270			
Ala	Val	Leu	Glu	Met	Leu	Glu	Glu	Ile	Ala	Ala	Asn	Gly	Gly	Asp	Ala
	275					280					285				
Thr	Asp	Phe	Met	Asn	Arg	Val	Lys	Asn	Lys	Glu	Lys	Gly	Val	Arg	Leu
	290				295				300						
Met	Gly	Phe	Gly	His	Arg	Val	Tyr	Lys	Asn	Tyr	Asp	Pro	Arg	Ala	Ala

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305		310		315		320
Ile Val Lys Asp Thr	Ala His Glu Ile	Leu Glu His Leu	Gly Gly Asp			
	325		330		335	
Pro Leu Leu Asp Leu	Ala Leu Lys Leu	Glu Glu Ile Ala	Leu Asn Asp			
	340		345		350	
Asp Tyr Phe Ile Ser	Arg Lys Leu Tyr	Pro Asn Val Asp	Phe Tyr Thr			
	355		360		365	
Gly Leu Ile Tyr Arg	Ala Met Gly Phe	Pro Thr Asp Phe	Phe Thr Val			
	370		375		380	
Leu Phe Ala Ile Gly	Arg Leu Pro Gly	Trp Ile Ala His	Tyr Arg Glu			
385		390		395		400
Gln Leu Ala Asp Pro	Gly Ala Lys Ile	Asn Arg Pro Arg	Gln Ile Tyr			
	405		410		415	
Thr Gly Glu Thr Ala	Arg Lys Ile Ile	Pro Arg Glu Glu	Arg			
	420		425		430	

&lt;210&gt; 91

&lt;211&gt; 1314

&lt;212&gt; DNA

&lt;213&gt; Brevibacterium lactofermentum

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1)..(1311)

&lt;400&gt; 91

atg ttt gaa agg gat atc gtg gct act gat aac aac aag gct gtc ctg	48
Met Phe Glu Arg Asp Ile Val Ala Thr Asp Asn Asn Lys Ala Val Leu	
1 5 10 15	
cac tac ccc ggt ggc gag ttc gaa atg gac atc atc gag gct tct gag	96
His Tyr Pro Gly Gly Glu Phe Glu Met Asp Ile Ile Glu Ala Ser Glu	
20 25 30	
ggt aac aac ggt gtt gtc ctg ggc aag atg ctg tct gag act gga ctg	144
Gly Asn Asn Gly Val Val Leu Gly Lys Met Leu Ser Glu Thr Gly Leu	
35 40 45	
atc act ttt gac cca ggt tat gtg agc act ggc tcc acc gag tgc aag	192
Ile Thr Phe Asp Pro Gly Tyr Val Ser Thr Gly Ser Thr Glu Ser Lys	
50 55 60	
atc acc tac atc gat ggc gat gcg gga atc ctg cgt tac cgc ggc tat	240
Ile Thr Tyr Ile Asp Gly Asp Ala Gly Ile Leu Arg Tyr Arg Gly Tyr	
65 70 75 80	
gac atc gct gat ctg gct gag aat gcc acc ttc aac gag gtt tct tac	288
Asp Ile Ala Asp Leu Ala Glu Asn Ala Thr Phe Asn Glu Val Ser Tyr	

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85					90					95							
cta	ctt	atc	aac	ggt	gaa	cta	cca	acc	cca	gat	gag	ctt	cac	aag	ttt	336	
Leu	Leu	Ile	Asn	Gly	Glu	Leu	Pro	Thr	Pro	Asp	Glu	Leu	His	Lys	Phe		
100					105					110							
aac	gac	gag	att	cgc	cac	cac	acc	ctt	ctg	gac	gag	gac	ttc	aag	tcc	384	
Asn	Asp	Glu	Ile	Arg	His	His	Thr	Leu	Leu	Asp	Glu	Asp	Phe	Lys	Ser		
115					120					125							
cag	ttc	aac	gtg	ttc	cca	cgc	gac	gct	cac	cca	atg	gca	acc	ttg	gct	432	
Gln	Phe	Asn	Val	Phe	Pro	Arg	Asp	Ala	His	Pro	Met	Ala	Thr	Leu	Ala		
130					135					140							
tcc	tcg	gtt	aac	att	ttg	tct	acc	tac	tac	cag	gat	cag	ctg	aac	cca	480	
Ser	Ser	Val	Asn	Ile	Leu	Ser	Thr	Tyr	Tyr	Gln	Asp	Gln	Leu	Asn	Pro		
145					150					155					160		
ctc	gat	gag	gca	cag	ctt	gat	aag	gca	acc	gtt	cgc	ctc	atg	gca	aag	528	
Leu	Asp	Glu	Ala	Gln	Leu	Asp	Lys	Ala	Thr	Val	Arg	Leu	Met	Ala	Lys		
165					170					175							
gtt	cca	atg	ctg	gct	gcg	tac	gca	cac	cgc	gca	cgc	aag	ggt	gct	ccf	576	
Val	Pro	Met	Leu	Ala	Ala	Tyr	Ala	His	Arg	Ala	Arg	Lys	Gly	Ala	Pro		
180					185					190							
tac	atg	tac	cca	gac	aac	tcc	ctc	aac	gcg	cgt	gag	aac	ttc	ctg	cgc	624	
Tyr	Met	Tyr	Pro	Asp	Asn	Ser	Leu	Asn	Ala	Arg	Glu	Asn	Phe	Leu	Arg		
195					200					205							
atg	atg	ttc	ggt	tac	cca	acc	gag	cca	tac	gag	atc	gac	cca	atc	atg	672	
Met	Met	Phe	Gly	Tyr	Pro	Thr	Glu	Pro	Tyr	Glu	Ile	Asp	Pro	Ile	Met		
210					215					220							
gtc	aag	gct	ctg	gac	aag	ctg	ctc	atc	ctg	cac	gct	gac	cac	gag	cag	720	
Val	Lys	Ala	Leu	Asp	Lys	Leu	Leu	Ile	Leu	His	Ala	Asp	His	Glu	Gln		
225					230					235					240		
aac	tgc	tcc	acc	tcc	acc	gtt	cgt	atg	atc	ggt	tcc	gca	cag	gcc	aac	768	
Asn	Cys	Ser	Thr	Ser	Thr	Val	Arg	Met	Ile	Gly	Ser	Ala	Gln	Ala	Asn		
245					250					255							
atg	ttt	gtc	tcc	atc	gct	ggt	ggc	atc	aac	gct	ctg	tcc	ggc	cca	ctg	816	
Met	Phe	Val	Ser	Ile	Ala	Gly	Gly	Ile	Asn	Ala	Leu	Ser	Gly	Pro	Leu		
260					265					270							
cac	ggt	ggc	gca	aac	cag	gct	gtt	ctg	gag	atg	ctc	gaa	gac	atc	aag	864	
His	Gly	Gly	Ala	Asn	Gln	Ala	Val	Leu	Glu	Met	Leu	Glu	Asp	Ile	Lys		
275					280					285							
aac	aac	cac	ggt	ggc	gac	gca	acc	gcg	ttc	atg	aac	aag	gtc	aag	aac	912	
Asn	Asn	His	Gly	Gly	Asp	Ala	Thr	Ala	Phe	Met	Asn	Lys	Val	Lys	Asn		
290					295					300							
aag	gaa	gac	ggc	gtc	cgc	ctc	atg	ggc	ttc	gga	cac	cgc	gtt	tac	aag	960	
Lys	Glu	Asp	Gly	Val	Arg	Leu	Met	Gly	Phe	Gly	His	Arg	Val	Tyr	Lys		
305					310					315					320		

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aac tac gat cca cgt gca gca atc gtc aag gag acc gca cac gag atc	1008
Asn Tyr Asp Pro Arg Ala Ala Ile Val Lys Glu Thr Ala His Glu Ile	
325 330 335	
ctc gag cac ctc ggt ggc gac gat ctt ctg gat ctg gca atc aag ctg	1056
Leu Glu His Leu Gly Gly Asp Asp Leu Leu Asp Leu Ala Ile Lys Leu	
340 345 350	
gaa gaa att gca ctg gct gat gat tac ttc atc tcc cgc aag ctc tac	1104
Glu Glu Ile Ala Leu Ala Asp Asp Tyr Phe Ile Ser Arg Lys Leu Tyr	
355 360 365	
ccg aac gta gac ttc tac acc ggc ctg atc tac cgc gca atg ggc ttc	1152
Pro Asn Val Asp Phe Tyr Thr Gly Leu Ile Tyr Arg Ala Met Gly Phe	
370 375 380	
cca act gac ttc ttc acc gta ttt gca atc ggt cgt ctg cca gga	1200
Pro Thr Asp Phe Phe Thr Val Leu Phe Ala Ile Gly Arg Leu Pro Gly	
385 390 395 400	
tgg atc gct cac tac cgc gag cag ctc ggt gca gca ggc aac aag atc	1248
Trp Ile Ala His Tyr Arg Glu Gln Leu Gly Ala Ala Gly Asn Lys Ile	
405 410 415	
aac cgc cca cgc cag gtc tac acc ggc aag gaa tcc cgc aag ttg gtt	1296
Asn Arg Pro Arg Gln Val Tyr Thr Gly Lys Glu Ser Arg Lys Leu Val	
420 425 430	
cct cgc gag gag cgc taa	1314
Pro Arg Glu Glu Arg	
435	

&lt;210&gt; 92

&lt;211&gt; 437

&lt;212&gt; PRT

&lt;213&gt; Brevibacterium lactofermentum

&lt;400&gt; 92

Met Phe Glu Arg Asp Ile Val Ala Thr Asp Asn Asn Lys Ala Val Leu	
1 5 10 15	
His Tyr Pro Gly Gly Glu Phe Glu Met Asp Ile Ile Glu Ala Ser Glu	
20 25 30	
Gly Asn Asn Gly Val Val Leu Gly Lys Met Leu Ser Glu Thr Gly Leu	
35 40 45	
Ile Thr Phe Asp Pro Gly Tyr Val Ser Thr Gly Ser Thr Glu Ser Lys	
50 55 60	
Ile Thr Tyr Ile Asp Gly Asp Ala Gly Ile Leu Arg Tyr Arg Gly Tyr	
65 70 75 80	
Asp Ile Ala Asp Leu Ala Glu Asn Ala Thr Phe Asn Glu Val Ser Tyr	
85 90 95	

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Leu	Leu	Ile	Asn	Gly	Glu	Leu	Pro	Thr	Pro	Asp	Glu	Leu	His	Lys	Phe	
			100						105					110		
Asn	Asp	Glu	Ile	Arg	His	His	Thr	Leu	Leu	Asp	Glu	Asp	Phe	Lys	Ser	
		115						120					125			
Gln	Phe	Asn	Val	Phe	Pro	Arg	Asp	Ala	His	Pro	Met	Ala	Thr	Leu	Ala	
	130						135					140				
Ser	Ser	Val	Asn	Ile	Leu	Ser	Thr	Tyr	Tyr	Gln	Asp	Gln	Leu	Asn	Pro	
145					150					155					160	
Leu	Asp	Glu	Ala	Gln	Leu	Asp	Lys	Ala	Thr	Val	Arg	Leu	Met	Ala	Lys	
			165						170						175	
Val	Pro	Met	Leu	Ala	Ala	Tyr	Ala	His	Arg	Ala	Arg	Lys	Gly	Ala	Pro	
			180					185						190		
Tyr	Met	Tyr	Pro	Asp	Asn	Ser	Leu	Asn	Ala	Arg	Glu	Asn	Phe	Leu	Arg	
	195						200					205				
Met	Met	Phe	Gly	Tyr	Pro	Thr	Glu	Pro	Tyr	Glu	Ile	Asp	Pro	Ile	Met	
	210					215					220					
Val	Lys	Ala	Leu	Asp	Lys	Leu	Leu	Ile	Leu	His	Ala	Asp	His	Glu	Gln	
225					230					235					240	
Asn	Cys	Ser	Thr	Ser	Thr	Val	Arg	Met	Ile	Gly	Ser	Ala	Gln	Ala	Asn	
			245					250						255		
Met	Phe	Val	Ser	Ile	Ala	Gly	Gly	Ile	Asn	Ala	Leu	Ser	Gly	Pro	Leu	
			260				265						270			
His	Gly	Gly	Ala	Asn	Gln	Ala	Val	Leu	Glu	Met	Leu	Glu	Asp	Ile	Lys	
	275						280					285				
Asn	Asn	His	Gly	Gly	Asp	Ala	Thr	Ala	Phe	Met	Asn	Lys	Val	Lys	Asn	
	290					295					300					
Lys	Glu	Asp	Gly	Val	Arg	Leu	Met	Gly	Phe	Gly	His	Arg	Val	Tyr	Lys	
305					310					315					320	
Asn	Tyr	Asp	Pro	Arg	Ala	Ala	Ile	Val	Lys	Glu	Thr	Ala	His	Glu	Ile	
			325						330					335		
Leu	Glu	His	Leu	Gly	Gly	Asp	Asp	Leu	Leu	Asp	Leu	Ala	Ile	Lys	Leu	
			340				345						350			
Glu	Glu	Ile	Ala	Leu	Ala	Asp	Asp	Tyr	Phe	Ile	Ser	Arg	Lys	Leu	Tyr	
	355						360					365				
Pro	Asn	Val	Asp	Phe	Tyr	Thr	Gly	Leu	Ile	Tyr	Arg	Ala	Met	Gly	Phe	
	370					375					380					
Pro	Thr	Asp	Phe	Phe	Thr	Val	Leu	Phe	Ala	Ile	Gly	Arg	Leu	Pro	Gly	
385					390					395					400	
Trp	Ile	Ala	His	Tyr	Arg	Glu	Gln	Leu	Gly	Ala	Ala	Gly	Asn	Lys	Ile	
			405						410					415		
Asn	Arg	Pro	Arg	Gln	Val	Tyr	Thr	Gly	Lys	Glu	Ser	Arg	Lys	Leu	Val	
			420					425					430			
Pro	Arg	G														

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435

<210> 93  
 <211> 1656  
 <212> DNA  
 <213> Corynebacterium thermoaminogenes

<220>  
 <221> CDS  
 <222> (309)..(1595)

&lt;400&gt; 93

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acgcccgaat cttcaacact atcgaagagg tcccaaccca cgcgttgacc cagggcttgg 60
gtactttgic ccgcgcgcaa aatatcgigt lggiggcaac tggccaagga aaagcagaca 120
gccatccgcg gaactgttga aggtccagtg actgcttctt gcccaggltc cattctgcaa 180
atgcacaaca acgccaccat catcgttgat gaagcagcag catccaagct gaaaaatgct 240
gaccattacc gtctcatgga gcaattaaag ctgcgctaga aacaaaaagg aaagtactgt 300
gtggggct atg cac aca gaa ctt tcc agt ttg cgc cct gcg tac cat gtg 350
      Met His Thr Glu Leu Ser Ser Leu Arg Pro Ala Tyr His Val
              1              5              10
act cct ccg cag ggc aga ctc aat gat ccc aat gga atg tac gtc gat 398
Thr Pro Pro Gln Gly Arg Leu Asn Asp Pro Asn Gly Met Tyr Val Asp
      15              20              25              30
gga gat acc ctc cac gtc tac tac cag cac gat cca ggt ttc ccc ttc 446
Gly Asp Thr Leu His Val Tyr Tyr Gln His Asp Pro Gly Phe Pro Phe
              35              40              45
gca cca aag cgc acc ggt tgg gct cac acc acc acg ccg ttg acc gga 494
Ala Pro Lys Arg Thr Gly Trp Ala His Thr Thr Thr Pro Leu Thr Gly
              50              55              60
ccg cag cga ttg cag tgg acg cac ctg ccc gat gct ctt tac ccg gat 542
Pro Gln Arg Leu Gln Trp Thr His Leu Pro Asp Ala Leu Tyr Pro Asp
              65              70              75
gta tcc tat gac ctg gat gga tgc tat tcc ggc gga gcc gta ttt tct 590
Val Ser Tyr Asp Leu Asp Gly Cys Tyr Ser Gly Gly Ala Val Phe Ser
              80              85              90
gac ggc acg ctt aaa ctt ttc tac acc ggc aac cga aaa att gac ggc 638
Asp Gly Thr Leu Lys Leu Phe Tyr Thr Gly Asn Arg Lys Ile Asp Gly
              95              100              105              110
aag cgc cgc gcc acc caa aac ctc gtc gaa gtc gag gac cca act ggg 686
Lys Arg Arg Ala Thr Gln Asn Leu Val Glu Val Glu Asp Pro Thr Gly
              115              120              125
ctg atg ggc ggc att cat cgc cgc tgc cct aaa aat ccg ctt atc gac 734
Leu Met Gly Gly Ile His Arg Arg Ser Pro Lys Asn Pro Leu Ile Asp

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			130			135			140						
gga	ccc	gcc	agc	ggt	ttt	acg	ccc	cac	tac	cgc	gat	ccc	atg	atc	agc
Gly	Pro	Ala	Ser	Gly	Phe	Thr	Pro	His	Tyr	Arg	Asp	Pro	Met	Ile	Ser
			145			150			155						
cct	gat	ggg	gat	ggt	tgg	aag	atg	gtt	ctt	ggg	gct	cag	cgc	gaa	aac
Pro	Asp	Gly	Asp	Gly	Trp	Lys	Met	Val	Leu	Gly	Ala	Gln	Arg	Glu	Asn
			160			165			170						
ctc	acc	ggt	gca	gcg	gtt	cta	tac	cgc	tcg	gca	gat	ctt	gaa	aac	tgg
Leu	Thr	Gly	Ala	Ala	Val	Leu	Tyr	Arg	Ser	Ala	Asp	Leu	Glu	Asn	Trp
			175			180			185			190			
gaa	ttc	tcc	ggt	gaa	atc	acc	ttt	gac	ctc	agc	gac	gca	caa	ccg	ggt
Glu	Phe	Ser	Gly	Glu	Ile	Thr	Phe	Asp	Leu	Ser	Asp	Ala	Gln	Pro	Gly
			195			200			205						
tct	gcc	cct	gat	ctc	gtt	cct	ggc	ggc	tac	atg	tgg	gaa	tgc	ccc	aac
Ser	Ala	Pro	Asp	Leu	Val	Pro	Gly	Gly	Tyr	Met	Trp	Glu	Cys	Pro	Asn
			210			215			220						
ctt	ttt	acg	ctt	cgc	gat	gaa	aaa	acc	ggc	gaa	gac	ctc	gat	gtg	ctg
Leu	Phe	Thr	Leu	Arg	Asp	Glu	Lys	Thr	Gly	Glu	Asp	Leu	Asp	Val	Leu
			225			230			235						
att	ttc	tgt	cca	caa	gga	tig	gac	cgt	atc	gat	gat	gag	gtt	act	cac
Ile	Phe	Cys	Pro	Gln	Gly	Leu	Asp	Arg	Ile	Asp	Asp	Glu	Val	Thr	His
			240			245			250						
tac	gca	agc	tct	gac	cag	tgc	gga	tat	gtc	gtc	ggc	aag	ctt	gaa	gaa
Tyr	Ala	Ser	Ser	Asp	Gln	Cys	Gly	Tyr	Val	Val	Gly	Lys	Leu	Glu	Glu
			255			260			265			270			
acg	acc	ttc	cgt	gtc	ctg	cga	gga	ttc	agc	gag	ctg	gat	ttc	ggt	cat
Thr	Thr	Phe	Arg	Val	Leu	Arg	Gly	Phe	Ser	Glu	Leu	Asp	Phe	Gly	His
			275			280			285						
gaa	ttc	tac	gcg	ccg	cag	gtt	gca	gtc	aac	ggt	tcc	gat	gcc	tgg	ctt
Glu	Phe	Tyr	Ala	Pro	Gln	Val	Ala	Val	Asn	Gly	Ser	Asp	Ala	Trp	Leu
			290			295			300						
gtg	ggc	tgg	atg	gga	tig	cct	gca	cag	gat	gat	cac	cca	aca	gtt	gcg
Val	Gly	Trp	Met	Gly	Leu	Pro	Ala	Gln	Asp	Asp	His	Pro	Thr	Val	Ala
			305			310			315						
cag	gaa	gga	tgg	gtg	cac	tgc	ctg	acc	gtt	cct	cgc	agg	ctt	cat	tig
Gln	Glu	Gly	Trp	Val	His	Cys	Leu	Thr	Val	Pro	Arg	Arg	Leu	His	Leu
			320			325			330						
cgt	aac	cat	gcg	atc	tat	caa	gag	ctt	ctt	ctc	cca	gaa	ggg	gag	tcg
Arg	Asn	His	Ala	Ile	Tyr	Gln	Glu	Leu	Leu	Leu	Pro	Glu	Gly	Glu	Ser
			335			340			345			350			
ggg	gta	act	aga	tct	gta	tta	ggt	tct	gaa	cct	gtc	cga	gta	gac	atc
Gly	Val	Thr	Arg	Ser	Val	Leu	Gly	Ser	Glu	Pro	Val	Arg	Val	Asp	Ile
			355			360			365						



118/123

cga gac aat gtt tcc ctc gag tgg gat ggt gtc cgg ttg tct gtg gat 1454  
 Arg Asp Asn Val Ser Leu Glu Trp Asp Gly Val Arg Leu Ser Val Asp  
 370 375 380  
 cgc gat ggc gat cgt cgt gta gct gaa gla aaa cct ggc gaa tta gtg 1502  
 Arg Asp Gly Asp Arg Arg Val Ala Glu Val Lys Pro Gly Glu Leu Val  
 385 390 395  
 atc gcg gac gat aat aca gcg att gag ata aca gca ggt cat ggc cag 1550  
 Ile Ala Asp Asp Asn Thr Ala Ile Glu Ile Thr Ala Gly His Gly Gln  
 400 405 410  
 gtt tcc ttc gct ttc cgc acc ttc aaa ggt gac act att gag aga 1595  
 Val Ser Phe Ala Phe Arg Thr Phe Lys Gly Asp Thr Ile Glu Arg  
 415 420 425  
 taagtcataa aaaagggcct tctgtggcgg attgtacaaa tacttcgcaa aatcccttga 1655  
 t 1656

&lt;210&gt; 94

&lt;211&gt; 429

&lt;212&gt; PRT

<213> *Corynebacterium thermoaminogenes*

&lt;400&gt; 94

Met His Thr Glu Leu Ser Ser Leu Arg Pro Ala Tyr His Val Thr Pro  
 1 5 10 15  
 Pro Gln Gly Arg Leu Asn Asp Pro Asn Gly Met Tyr Val Asp Gly Asp  
 20 25 30  
 Thr Leu His Val Tyr Tyr Gln His Asp Pro Gly Phe Pro Phe Ala Pro  
 35 40 45  
 Lys Arg Thr Gly Trp Ala His Thr Thr Thr Pro Leu Thr Gly Pro Gln  
 50 55 60  
 Arg Leu Gln Trp Thr His Leu Pro Asp Ala Leu Tyr Pro Asp Val Ser  
 65 70 75 80  
 Tyr Asp Leu Asp Gly Cys Tyr Ser Gly Gly Ala Val Phe Ser Asp Gly  
 85 90 95  
 Thr Leu Lys Leu Phe Tyr Thr Gly Asn Arg Lys Ile Asp Gly Lys Arg  
 100 105 110  
 Arg Ala Thr Gln Asn Leu Val Glu Val Glu Asp Pro Thr Gly Leu Met  
 115 120 125  
 Gly Gly Ile His Arg Arg Ser Pro Lys Asn Pro Leu Ile Asp Gly Pro  
 130 135 140  
 Ala Ser Gly Phe Thr Pro His Tyr Arg Asp Pro Met Ile Ser Pro Asp  
 145 150 155 160  
 Gly Asp Gly Trp Lys Met Val Leu Gly Ala Gln Arg Glu Asn Leu Thr  
 165 170 175

119/123

Gly	Ala	Ala	Val	Leu	Tyr	Arg	Ser	Ala	Asp	Leu	Glu	Asn	Trp	Glu	Phe
			180					185					190		
Ser	Gly	Glu	Ile	Thr	Phe	Asp	Leu	Ser	Asp	Ala	Gln	Pro	Gly	Ser	Ala
		195					200					205			
Pro	Asp	Leu	Val	Pro	Gly	Gly	Tyr	Met	Trp	Glu	Cys	Pro	Asn	Leu	Phe
		210				215					220				
Thr	Leu	Arg	Asp	Glu	Lys	Thr	Gly	Glu	Asp	Leu	Asp	Val	Leu	Ile	Phe
225					230					235					240
Cys	Pro	Gln	Gly	Leu	Asp	Arg	Ile	Asp	Asp	Glu	Val	Thr	His	Tyr	Ala
			245					250						255	
Ser	Ser	Asp	Gln	Cys	Gly	Tyr	Val	Val	Gly	Lys	Leu	Glu	Glu	Thr	Thr
		260						265					270		
Phe	Arg	Val	Leu	Arg	Gly	Phe	Ser	Glu	Leu	Asp	Phe	Gly	His	Glu	Phe
	275						280					285			
Tyr	Ala	Pro	Gln	Val	Ala	Val	Asn	Gly	Ser	Asp	Ala	Trp	Leu	Val	Gly
	290					295					300				
Trp	Met	Gly	Leu	Pro	Ala	Gln	Asp	Asp	His	Pro	Thr	Val	Ala	Gln	Glu
305					310					315					320
Gly	Trp	Val	His	Cys	Leu	Thr	Val	Pro	Arg	Arg	Leu	His	Leu	Arg	Asn
			325					330						335	
His	Ala	Ile	Tyr	Gln	Glu	Leu	Leu	Leu	Pro	Glu	Gly	Glu	Ser	Gly	Val
		340						345					350		
Thr	Arg	Ser	Val	Leu	Gly	Ser	Glu	Pro	Val	Arg	Val	Asp	Ile	Arg	Asp
		355					360					365			
Asn	Val	Ser	Leu	Glu	Trp	Asp	Gly	Val	Arg	Leu	Ser	Val	Asp	Arg	Asp
	370					375					380				
Gly	Asp	Arg	Arg	Val	Ala	Glu	Val	Lys	Pro	Gly	Glu	Leu	Val	Ile	Ala
385					390					395					400
Asp	Asp	Asn	Thr	Ala	Ile	Glu	Ile	Thr	Ala	Gly	His	Gly	Gln	Val	Ser
			405					410						415	
Phe	Ala	Phe	Arg	Thr	Phe	Lys	Gly	Asp	Thr	Ile	Glu	Arg			
		420						425							

&lt;210&gt; 95

&lt;211&gt; 35

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence: primer for  
amplifying scrB gene

120/123

<400> 95  
glacataattg tcgttagaac gcgtaatacg actca 35

<210> 96  
<211> 35  
<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:primer for  
amplifying scrB gene

<400> 96  
cgtagaacg cgtaatacga ctactatag ggaga 35

<210> 97  
<211> 30  
<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:primer for  
LA cloning of scrB

<400> 97  
gtaaagagcg tcgggcaggt gcgtccactg 30

<210> 98  
<211> 30  
<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:primer for  
LA cloning of scrB

<400> 98  
gggtgtgagcc cagccgggtgc gctttgggtgc 30

<210> 99

121/123

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
LA cloning of scrB

&lt;400&gt; 99

atcagccctg atggatgatgg ttggaaaatg

30

&lt;210&gt; 100

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
LA cloning of scrB gene

&lt;400&gt; 100

ggtgcagcgg ttctataccg ctgcacagat

30

&lt;210&gt; 101

&lt;211&gt; 32

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying scrB gene

&lt;400&gt; 101

ggcccgggac gcccgattct tcaacactat cg

32

&lt;210&gt; 102

&lt;211&gt; 32

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

122/123

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying scrB gene

&lt;400&gt; 102

ggccccgggga tcaagggatt ttgcgaagta tt

32

&lt;210&gt; 103

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying icd gene

&lt;400&gt; 103

gaagatctct atgaccagcg catcaagctg

30

&lt;210&gt; 104

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying icd gene

&lt;400&gt; 104

gaagatctgg tcatcccaga acctgatcac

30

&lt;210&gt; 105

&lt;211&gt; 32

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gdh gene

123/123

&lt;400&gt; 105

gcgcctgcag gtccgagggt gtgcgttcgg ca

32

&lt;210&gt; 106

&lt;211&gt; 32

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gdh gene

&lt;400&gt; 106

gcgcctgcag gcaccaggat gccctcaacc ag

32

&lt;210&gt; 107

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 107

ggggtaccga tcactataac cccacagcac

30

&lt;210&gt; 108

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Description of Artificial Sequence:primer for  
amplifying gltA gene

&lt;400&gt; 108

ggggtaccct ggctgatctg aactaggcgc

30